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Description

[0001] This invention relates to a female-sterile plant and to its reproductive material (e.g., seeds), in which the cells have been transformed so that a foreign DNA sequence is stably integrated into their nuclear genome. The foreign DNA sequence of this invention contains a first foreign DNA (hereinafter the "female-sterility DNA") that: 1) encodes a first RNA, protein or polypeptide which, when produced or overproduced in a cell of a flower, particularly a female organ thereof, or a seed or an embryo of a plant, disturbs significantly the metabolism, functioning and/or development of the cell of the flower or seed or embryo; and 2) is in the same transcriptional unit as, and under the control of, a first promoter which is capable of directing expression of the female-sterility DNA selectively in cells of the flowers, particularly one or more of their female organs, or seeds or embryos of the plant. In particular, this invention relates to such a nuclear female-sterile plant and its reproductive material, in which the foreign DNA sequence of this invention is a foreign chimaeric DNA sequence that can also contain a second foreign DNA (the "marker DNA") that: encodes a second RNA, protein or polypeptide which, when present at least in a specific tissue or specific cells of the plant, renders the entire plant easily separable from other plants that do not contain the second RNA, protein or polypeptide at least in the specific tissue or specific cells; 2) is in the same transcriptional unit as, and under the control of, a second promoter which is capable of directing expression of the marker DNA in at least the specific tissue or the specific cells of the plant; and 3) is in the same genetic locus of the nuclear genome of the cells of the plant as the female-sterility DNA.

[0002] This invention also relates to a foreign chimaeric DNA sequence that contains at least one female-sterility DNA under the control of at least one first promoter and that can also contain, adjacent to the female-sterility DNA(s) and the first promoter(s), at least one marker DNA under the control of at least one second promoter.

[0003] This invention further relates to a vector that contains the foreign DNA sequence of this invention and is suitable for the transformation of a plant cell, whereby the foreign DNA sequence is stably integrated into the nuclear genome of the cell.

[0004] This invention still further relates to cells of a plant and to plant cell cultures, the nuclear genomes of which are transformed with the foreign DNA sequence.

[0005] This invention yet further relates to a process for producing a nuclear female-sterile male-fertile plant and its reproductive material containing the foreign DNA sequence in which the female-sterility DNA: 1) is under the control of the first promoter; 2) is stably integrated into the nuclear genome of the plant's cells; 3) can be expressed selectively in cells of each flower, particularly a female organ thereof, or each seed or each embryo of the plant in the form of the first RNA, protein or polypeptide; and optionally 4) is in the same genetic locus as the marker DNA under the control of the second promoter.

[0006] The invention further relates to a process for producing hybrid seeds, which grow into hybrid plants, by crossing: 1) the female-sterile plant of this invention which may include, in its nuclear genome, the marker DNA, preferably encoding a protein conferring a resistance to a herbicide on the plant; and 2) a female-fertile plant without the marker DNA in its genome. This invention particularly relates to such a process for producing hybrid seeds on a commercial scale, preferably in a substantially random population, without the need for extensive hand-labor.

Background of the invention

[0007] Hybridization of plants is recognized as an important process for producing offspring having a combination of the desirable traits of the parent plants. The resulting hybrid offspring often has the ability to outperform the parents in different traits, such as in yield, adaptability to environmental changes, and disease resistance. This ability is called "heterosis" or "hybrid vigor". As a result, hybridization has been used extensively for improving major crops, such as corn, sugar beet and sunflower. For a number of reasons, primarily related to the fact that most plants are capable of undergoing both self-pollination and cross-pollination, the controlled cross-pollination of plants without significant self-pollination, to produce a harvest of hybrid seeds, has been difficult to achieve on a commercial scale.

[0008] In nature, the vast majority of crop plants produce male and female reproductive organs on the same plant, usually in close proximity to one another in the same flower. This favors self-pollination. Some plants, however, are exceptions as a result of the particular morphology of their reproductive organs which favors cross-pollination. These plants produce hybrid offspring with improved vigor and adaptability. One such morphology in *Cannabis ssp.* (hemp) involves male and female reproductive organs on separate plants. Another such morphology in *Zea mays* (corn) involves male and female reproductive organs on different parts of the same plant. Another such morphology in *Elaeis guineensis* (oil palm) involves male and fertile female gametes which become fertile at different times in the plant's development.

[0009] Some other plant species, such as *Ananas comosus* (pineapple), favor cross-pollination through the particular physiology of their reproductive organs. Such plants have developed a so-called "self-incompatibility system" whereby the pollen of one plant is not able to fertilize the female gamete of the same plant or of another plant with the same genotype.

[0010] Some other plant species favor cross-pollination by naturally displaying the so-called genomic characteristic of "male sterility". By this characteristic, the plants' anthers degenerate before pollen, produced by the anthers, reaches maturity. See: "Male-sterility in Higher Plants", M.L.H. Kaul, 1987, in: Monographs on Theoretical and Applied Genetics 10, Edit. Springer Verlag. Such a natural male-sterility characteristic is believed to result from a wide range of natural mutations, most often involving recessive deficiencies, and this characteristic can not easily be maintained in plant species that predominantly self-pollinate, since under natural conditions, no seeds will be produced.

[0011] Some other plants favor cross-pollination by naturally displaying the character of "female-sterility" due to a deficient functioning of either the female gametophyte, the female gamete, the female zygote, or the seed. These plants produce no viable seeds. There are many different mutations that can lead to this condition, involving all stages of development of a specific tissue of the female reproductive organ. This characteristic distinguishes female-sterility from the more widely known phenomena of male-sterility and self-incompatibility. Although reducing the number of offspring a species can produce, the female-sterility trait has some evolutionary advantages in nature for some plants, especially for perennials. In perennials, the rate of vegetative growth is to a large extent determined by the distribution of biomass between vegetative and reproductive plant tissues. Female-sterile plants therefore tend to grow more vigorously than the female-fertile plants.

[0012] Although female-sterility inducing mutations probably occur as frequently as male-sterility inducing mutations, female-sterility inducing mutations are much less used in plant breeding and seed production and consequently much less studied, and only few examples of such mutations exist.

[0013] A well documented illustration of natural female-sterility is the oil palm (*Elaeis guineensis*) where the so-called "pisifera" condition is characterized by the inability of the developing seed to produce a shell. As a result, the developing seed aborts in an early stage, and no ripe fruit is formed. The gene encoding the "pisifera" genotype acts as a semi-dominant allele. Plants homozygous for the allele are not capable of producing a seed shell and consequently no ripe fruit or seeds. Plants heterozygous for the allele produce ripe fruit and seeds with a thin shell (0.5 to 2 mm), while wild-type plants (which do not carry the allele) produce ripe fruit and seeds with shells of 2 to 6 mm thickness. These two genotypes are indistinguishable in seed yield, and their genotype is determined by that of the female parent plant. In oil palm breeding, the "pisifera" type is used as the male parent plant in all commercial seed production. By crossing pollen from the "pisifera" palms with the wild-type female parent plants, a homogeneous F1 hybrid seed population, producing thin-shelled fruit, is obtained.

[0014] Another example of a plant with a natural female-sterility, used for the commercial production of hybrid seed, is alfalfa. Alfalfa was known to have male-sterility genes, but in testing a hybrid seed production system in which male-sterile and male fertile plants were sown in separate bands, it appeared that a negligible amount of hybrid seeds was produced. This low production was due to the fact that honeybees, responsible for pollination, have low affinity for male-sterile plants, favoring the self-pollination of the male-fertile plants. To obtain good seed set, it seemed necessary to interplant very closely to each other (thus not in separate rows) the male-fertile and the male-sterile parent plants. This was made possible when a female-sterility gene was discovered and bred into the male-fertile plants. Consequently, the only seeds which could be produced in the randomly sown plots were hybrid seeds obtained by cross-pollination between the female-sterile and the male-sterile parent plants.

[0015] For other crops, female-sterility has been reported, such as sorghum (Casady et al (1960) J. Hered. 51, 35-38), cotton (Tustus and Meyer (1963) J. Hered. 54, 167-168), tomato (Honma and Pratak (1964) J. Hered. 55, 143-145), wheat (Gotzov and Dzelepov (1974) Gen. Plant Breed. Z, 480-487), and pearl millet (Hanna and Powel (1974) J. Hered. 65, 247-249). There are, however, several problems in maintaining the female-sterile lines, and for this reason, such lines are not used on a commercially important scale.

[0016] Compared with male-sterility, the use of female-sterility offers some other advantages in the production of hybrid seeds. Female-sterility allows the production of fruits without seeds and enhanced vegetative biomass production and can, in some cases, induce more flower-setting within one season.

Summary of the Invention

[0017] In accordance with this invention a DNA, such as a nuclear DNA of a cell of a plant, is provided which contains a first chimeric DNA which comprises:

- (a) a female sterility DNA encoding a first RNA, protein or polypeptide, capable, when produced in cells of a female reproductive organ of the plant, of disturbing significantly the metabolism, functioning and/or development of the female reproductive organ; and
- (b) a first promoter capable of directing expression of this DNA, selectively in cells of the female reproductive organs of the plant, the female sterility DNA being in the same transcriptional unit as, and under the control of the first promoter.

The first promoter preferably directs expression especially in style cells, stigma cells, ovary cells, ovule cells and/or septum cells, particularly in stigma and/or style cells. The female-sterility DNA preferably encodes a ribonuclease, such as barnase.

[0018] The DNA of this invention can also contain a second chimeric DNA comprising:

(c) a marker DNA encoding a second RNA, protein or polypeptide which, when present at least in a specific tissue or in at least specific cells of a plant, renders the plant easily separable from other plants which do not contain the second RNA, protein or polypeptide in the specific tissue or specific cells; and

(d) a second promoter capable of directing expression of the marker DNA at least in the specific tissue or specific cells; the marker DNA being in the same transcriptional unit as, and under the control of, the second promoter.

The marker DNA is preferably a herbicide resistance gene.

[0019] Also in accordance with this invention is provided a DNA comprising the female-sterility DNA under the control of a first promoter, and that may also comprise the marker DNA and the second promoter, as well as at least one additional DNA encoding: a transit peptide capable of transporting the first protein or polypeptide and/or the second protein or polypeptide into a chloroplast or mitochondrion of a plant cell; and/or a secretory signal peptide capable of secreting the first protein or polypeptide, and/or the second protein or polypeptide out of a plant cell or plant tissue.

[0020] Further in accordance with this invention are provided; a cell of a plant, a plant cell culture, a plant, a plant seed, and a seedless fruit containing the DNA of this invention. Also provided are a pair of parent plants comprising a) a female-sterile parent plant containing the DNA of this invention, and b) a male-fertile parent plant.

[0021] In particular this invention provides a female-sterile plant containing a foreign DNA incorporated in the nuclear genome of its cells, wherein the foreign DNA comprises :

(a) a female-sterility DNA encoding a first RNA, protein or polypeptide, capable when produced in cells of a female reproductive organ of the plant, of disturbing significantly the metabolism, functioning and/or development of the female reproductive organ; and

(b) a first promoter capable of directing expression of this DNA selective in cells of the female reproductive organs of the plant, the female sterility DNA being in the same transcriptional unit as, and under the control of the first promoter,

provided that, if the first promoter is a promoter capable of directing expression of the female-sterility DNA selectively in female gametes, or in cells derived from female gametes, the nuclear genome of the plant is homozygous.

[0022] Still further in accordance with this invention a process is provided for producing a female-sterile plant and reproductive material, e.g. seeds, thereof, by transforming a cell of a plant with a DNA comprising the female-sterility DNA under control of the first promoter, regenerating the female-sterile plant, and optionally, obtaining the reproduction material or progeny of the female-sterile plant.

[0023] Still further in accordance with this invention a process for producing a seed is provided which comprises the steps of:

(a) providing a female-sterile plant which contains a DNA comprising a female-sterility DNA under the control of a first promoter and a marker DNA, which is a gene conferring resistance to a herbicide, under the control of a second promoter,

(b) cross-pollinating i) the female-sterile plant, and ii) a female-fertile plant, preferably a male-sterile plant, without the marker DNA and/or the second promoter, and

(c) recovering the seed from the female-fertile plant,

wherein the process comprises removing undesired female-fertile plants by application of the herbicide.

Detailed Description of the Invention

[0024] In accordance with this invention, a female-sterile male-fertile plant is produced from a single cell of a plant by transforming the plant cell in a well known manner to stably insert, into the nuclear genome of the cell, the foreign DNA sequence of this invention. The foreign DNA sequence comprises at least one female-sterility DNA that is under the control of, and fused at its 5' end to, the first promoter and is fused at its 3' end to suitable transcription termination (or regulation) signals, including a polyadenylation signal. Thereby, the first RNA, protein or polypeptide is produced or overproduced selectively in cells of all the flowers, particularly in one or more female organs thereof, and/or in all the seeds and/or in all the embryos of the plant so as to render the plant female-sterile. The foreign DNA sequence can also comprise at least one marker DNA that is under the control of, and is fused at its 5' end to, the second promoter

and is fused at its 3' end to suitable transcription termination signals, including a polyadenylation signal. The marker DNA is preferably in the same genetic locus as the female-sterility DNA, whereby the second RNA, protein or polypeptide is produced in at least the specific tissue or specific cells of the female-sterile plant so that the plant can be easily distinguished and/or separated from other plants that do not contain the second RNA, protein or polypeptide in the specific tissue or specific cells. This guarantees, with a high degree of certainty, the joint segregation of both the female-sterility DNA and the marker DNA into offspring of the plant.

[0025] This invention relates to any plant, the nuclear genome of which can be transformed with a female-sterility DNA under the control of a first promoter that can direct expression of the female-sterility DNA selectively in the cells of the flowers, particularly a female organ thereof, whereby the plant can be both self-pollinated or cross-pollinated.

[0026] The cell of the plant (particularly a plant capable of being infected with *Agrobacterium*) is preferably transformed in accordance with this invention, using a vector that is a disarmed Ti-plasmid containing the foreign DNA sequence and carried by *Agrobacterium*. This transformation can be carried out using procedures described, for example, in European patent publications 0,116,718 and 0,270,822. Preferred Ti-plasmid vectors contain the foreign DNA sequence between the border sequences, or at least located to the left of the right border sequence, of the T-DNA of the Ti-plasmid. Of course, other types of vectors can be used to transform the plant cell, using procedures such as direct gene transfer (as described, for example, in European patent publication 0,223,247), pollen mediated transformation (as described, for example, in European patent publication 0,270,356, PCT publication WO85/01856, and European patent publication 0,275,069), *in vitro* protoplast transformation (as described, for example, in US patent 4,684,611), plant RNA virus-mediated transformation (as described, for example, in European patent publication 0,067,553 and US patent 4,407,956) and liposome-mediated transformation (as described, for example, in US patent 4,536,475).

[0027] Preferably, a nuclear female-sterile male-fertile plant of this invention is provided by transforming a plant cell with a disarmed Ti-plasmid vector containing the foreign DNA sequence with a female-sterility DNA under the control of a first promoter and preferably a marker DNA under the control of a second promoter. The marker DNA can be upstream or downstream of the female-sterility DNA in the Ti-plasmid vector, but preferably, the two are adjacent to one another and are located between the border sequences or at least located to the left of the right border sequence of the Ti-plasmid vector, so that they are properly transferred together into the nuclear genome of the plant cell. However, if desired, the cell can initially be transformed with a foreign DNA sequence containing a female-sterility DNA and a first promoter and can subsequently be transformed with a marker DNA and a second promoter, inserted into or near the genetic locus of the female-sterility DNA in the cell's nuclear genome, or this transformation can be carried out vice versa. Suitable vectors for this purpose are the same as those discussed above for transforming cells with the foreign DNA sequence. The preferred vector is a disarmed Ti-plasmid vector.

[0028] The selection of the female-sterility DNA of this invention is not critical. A suitable female-sterility DNA can be selected and isolated in a well-known manner, so that it encodes the first RNA, protein or polypeptide which significantly disturbs adversely the proper metabolism and/or functioning and/or development of any cell of a flower and/or seed and/or embryo in which the female-sterility DNA is expressed, preferably leading thereby to the death of such cell. Preferred examples of female-sterility DNAs encode: RNases such as RNase T1 (which degrades RNA molecules by hydrolyzing the bond after any guanine residue) and Barnase; DNases such as an endonuclease (e.g., *EcoRI*); or proteases such as a papain (e.g., papain zymogen and papain active protein). Other preferred examples of female-sterility DNAs encode: ribonucleases such as T₂ (Kawata et al (1988) Eur. J. Biochem. 176, 683-697) or Rh (Horiuchi et al (1988) J. Biochem. 103, 408-418); or glycoproteins such as are encoded by the S1, S2, S3, S6 and S7 alleles, particularly of *Nicotiana glauca* (McClure et al (1989) Nature 342, 955-957).

[0029] Other examples of female-sterility DNAs encode enzymes which catalyze the synthesis of phytohormones, such as: isopentenyl transferase which is an enzyme that catalyzes the first step in cytokinin biosynthesis and is encoded by gene 4 of *Agrobacterium* T-DNA; or one or both of the enzymes involved in the synthesis of auxin and encoded by gene 1 and gene 2 of *Agrobacterium* T-DNA. Yet other examples of female-sterility DNAs encode: glucanases; lipases such as phospholipase A₂ (Verheij et al (1981) Rev. Biochem. Pharmacol. 91, 92-203); lipid peroxidases; or plant cell wall inhibitors. Still other examples of female-sterility DNAs encode proteins toxic to plants cells, such as a bacterial toxin (e.g., the A-fragment of diphtheria toxin or botulin).

[0030] Still another example of a female-sterility DNA is an antisense DNA: i) which encodes a strand of RNA complementary to a strand of RNA that is endogenous to, and naturally transcribed in, the cells of the flower, seed or embryo of the plant of this invention and ii) which is under the control of an endogenous promoter as described, for example, in European patent publication 0,223,399. Such an antisense DNA can be transcribed into an RNA sequence capable of binding to the coding and/or non-coding portion of an RNA, naturally produced in the cell of the flower, seed or embryo, so as to inhibit the translation of the naturally produced RNA. Examples of such an antisense DNA are the antisense DNAs of: the STMG-type genes, such as the STMG07 gene, the STMG08 gene, the STMG4B12 gene and the STMG3C9 gene of Example 2 herein; the KT13 gene (Jofuku and Goldberg (1989) The Plant Cell 1, 1079-1093); a gene encoding a seed-specific storage protein, such as a 2S albumin (Krebbers et al (1988) Plant Physiol. 87, 859-866;

Altenbach et al (1987) *Plant Molecular Biol.* **8**, 239-250; (Scolfield and Crouch (1987) *J. Biol. Chem.* **262**, 12202-12208); or a gene corresponding to cDNA clone pMON9608 (Gasser et al (1989) *The Plant Cell* **1**, 15). Such antisense DNAs can be naturally expressed in flower, seed or embryo cells of the plant under the control of the endogenous promoter of the complementary endogenous DNA strand (or gene) of the plant, for example: in the style (with the antisense DNA of the STMG07, STMG08, STMG4B12 or STMG3C9 gene); in the embryo axis (with the antisense DNA of the KT13 gene); in seeds (with the antisense DNA of a 2S albumin-encoding gene); and in ovule cells (with the antisense DNA of PMON9608).

[0031] A further example of a female-sterility DNA encodes a specific RNA enzyme (i.e., a so-called "ribozyme"), capable of highly specific cleavage against a given target sequence as described by Haseloff and Gerlach (1988) *Nature* **334**, 585-591. Such a ribozyme is, for example, the ribozyme targeted against the RNA encoded by the STMG07 gene, the STMG08 gene, the STMG4B12 gene, the STMG3C9 gene, the KT13 gene, a gene encoding a seed-specific storage protein such as a 2S albumin or the gene corresponding to cDNA clone pMON9608.

[0032] Still other examples of female-sterility DNAs encode products which can render the flowers, seeds and/or embryos susceptible to a specific disease, such as a fungus infection. Such a female-sterility DNA can be used in a plant, in which all other cells or tissues, in which the female-sterility DNA is not expressed, are resistant to the specific disease.

[0033] Yet another example of a female-sterility DNA comprises a combination of: 1) a first gene encoding a viral dependent RNA polymerase, such as TNV replicase (Meulewater et al (1990) *Virology* **177**, 1-11), under the control of a first promoter of this invention; and 2) a negative strand (i.e., antisense DNA) of a second gene: i) which encodes a first protein or polypeptide of this invention that, when produced or overproduced in the plant cell of this invention, disturbs significantly cell metabolism, development and/or functioning, ii) which is fused at its 3' end to a viral RNA replication recognition sequence or so-called "viral subgenomic promoter", such as the TNV subgenomic promoter (Lexis et al (1990) *J. of Virology* **64** (4), 1726-1733); and iii) which is fused at its 5' end to, and under the control of, another suitable promoter, such as a first promoter of this invention, capable of directing expression of the negative strand in the plant cell of this invention. The viral subgenomic promoter sequence is specifically recognized by the viral dependent RNA polymerase encoded by the first gene. This recognition leads to the repeated replication of the negative strand of the second gene as a sense strand, which leads to the synthesis of the first protein or polypeptide. Both the first gene and the negative strand are provided in the nuclear genome of the plant cell of this invention. This can be achieved by one transformation event, by two consecutive transformation events, or by crossing a plant having the first gene inserted into its genome with a plant having the negative strand inserted into its genome.

[0034] By "foreign" with regard to the foreign DNA sequence of this invention is meant that the foreign DNA sequence contains a foreign female-sterility DNA and/or a foreign first promoter. By "foreign" with regard to a DNA, such as a female-sterility DNA and a first promoter, as well as a marker DNA, a second promoter and any other DNA in the foreign DNA sequence, is meant that such a DNA is not in the same genomic environment in a plant cell, transformed with such a DNA in accordance with this invention, as is such a DNA when it is naturally found in the cell of the plant, bacteria, animal, fungus, virus, or the like, from which such a DNA originates. This means, for example, that a foreign female-sterility DNA or marker DNA can be: 1) a nuclear DNA in a plant of origin; 2) endogenous to the transformed plant cell (i.e., from a plant of origin with the same genotype as the plant being transformed); and 3) within the same transcriptional unit as its own endogenous promoter and 3' end transcription regulation signals (from the plant of origin) in the foreign DNA sequence of this invention in the transformed plant cell; but 4) inserted in a different place in the nuclear genome of the transformed plant cell than it was in the plant of origin so that it is not surrounded in the transformed plant cell by the genes which surrounded it naturally in the plant of origin. A foreign female-sterility or marker DNA can also, for example, be: 1) a nuclear DNA in a plant of origin; and 2) endogenous to the transformed plant cell; but 3) in the same transcriptional unit as a different (i.e., not its own) endogenous promoter and/or 3' end transcription regulation signals in a foreign chimaeric DNA sequence of this invention in a transformed plant cell. A foreign female-sterility or marker DNA can also, for example, be: 1) a nuclear DNA in a plant of origin; and 2) endogenous to the transformed plant cell; but 3) in the same transcriptional unit as a heterologous promoter and/or 3' end transcription regulation signals in a foreign chimaeric DNA sequence of this invention in a transformed plant cell. A foreign female-sterility or marker DNA can also, for example, be heterologous to the transformed plant cell and in the same transcriptional unit as an endogenous promoter and/or 3' transcription regulation signals (e.g., from the nuclear genome of a plant with the same genotype as the plant being transformed) in a foreign chimaeric DNA sequence of this invention in a transformed plant cell. An example of a foreign female-sterility DNA could come from the nuclear genome of a plant with the same genotype as the plant being transformed and encode a catalytic enzyme, such as a protease or ribonuclease, that is endogenous to cells of the flowers, seeds and/or embryos of the plant being transformed, so that the enzyme is overproduced in transformed cells of the flowers, seeds and/or embryos in order to disturb significantly their metabolism, functioning and/or development. Preferably, the female-sterility DNA and the marker DNA are each heterologous to the plant cell being transformed.

[0035] By "heterologous" with regard to a DNA, such as a female-sterility DNA, a first promoter, a marker DNA, a

second promoter and any other DNA in the foreign DNA sequence of this invention, is meant that such a DNA is not naturally found in the nuclear genome of cells of a plant with the same genotype as the plant being transformed. Examples of heterologous DNAs include chloroplast and mitochondrial DNAs obtained from a plant with the same genotype as the plant being transformed, but preferred examples are chloroplast, mitochondrial, and nuclear DNAs from plants having a different genotype than the plant being transformed, DNAs from animal and bacterial genomes, and chromosomal and plasmidial DNAs from fungal and viral genomes.

[0036] By "chimaeric" with regard to the foreign DNA sequence of this invention is meant that at least one of its female-sterility DNAs: 1) is not naturally found under the control of its first promoter for the one female-sterility DNA; and/or 2) is not naturally found in the same genetic locus as at least one of its marker DNAs. Examples of foreign chimaeric DNA sequences of this invention comprise: a female-sterility DNA of bacterial origin under the control of a first promoter of plant origin; and a female-sterility DNA of plant origin under the control of a first promoter of plant origin and in the same genetic locus as a marker DNA of bacterial origin.

[0037] By "flower" is meant to include the entire organ of a flower, as well as one or more of its individual parts such as its shoot axis, sepals, petals, male reproductive organs (or stamens) and female reproductive organs (or carpels), whose wholly or partly, retarded or arrested development in accordance with this invention prevents the development of viable seeds in the flower but not the development and propagation of its male gametes; by "female organ" is meant the entire organ of a flower that is involved in the production of female gametes and/or viable seeds and/or viable embryos, as well as one or more of its individual parts such as its ovule, ovary, style, stigma, corolla, disc, septum, calyx and placenta tissue. By "embryo" is meant to include the entire embryo of a plant, as well as one or more of its individual parts such as its embryo axis and embryo cotyledons.

[0038] So that the female-sterility DNA of this invention is expressed selectively in cells of the flowers, particularly one or more of their female organs, in cells of the seeds and/or in cells of the embryos of the plants of this invention, it is preferred that the first promoter, which controls the female-sterility DNA in the foreign DNA sequence, be a promoter capable of directing gene expression selectively in cells of the flowers, seeds and/or embryos of the plant. Such a flower-, seed- and/or embryo-specific promoter can be an endogenous promoter or an exogenous promoter and can be from the nuclear genome or from the mitochondrial or chloroplast genome of a plant cell. In any event, the first promoter is foreign to the nuclear genome of the plant cell, being transformed. Preferably, the first promoter causes the female-sterility DNA to be expressed only in cells of one or more specific tissues of the flowers, preferably one or more female organs thereof, or of the seeds or of the embryos, especially in style cells, ovary cells, septum cells, seedcoat cells, endosperm cells, embryo axis cells and/or embryo cotyledon cells.

[0039] The first promoter of this invention can be selected and isolated in a well known manner from a plant, to be rendered female-sterile, so that the first promoter directs expression of the female-sterility DNA selectively in cells of the flowers, seeds and/or embryos of the plant, so as to kill or disable the plant's flowers, seeds and/or embryos and render the plant incapable of producing fertile female gametes, viable seeds and/or viable embryos. The first promoter is preferably also selected and isolated so that it is effective to prevent expression of the female-sterility DNA in other parts of the plant that are not involved in the production of fertile female gametes, viable seeds and/or viable embryos, especially in male organs of the flowers, so that the plant remains male-fertile. For example, a suitable flower-specific (preferably female reproductive organ-specific), seed-specific or embryo-specific first promoter can be identified and isolated in a plant, to be made female-sterile, by:

1. searching for an mRNA which is only present in the plant during the development of its flowers, seeds or embryos, preferably its ovary, style, placenta, calyx, scutellum, septum, seedcoat, endosperm or embryo cotyledons;
2. isolating this flower-, seed- or embryo-specific mRNA;
3. preparing a cDNA from this specific mRNA;
4. using this cDNA as a probe to identify the regions in the plant genome which contain DNA coding for this specific mRNA; and then
5. identifying the portion of the plant genome that is upstream (i.e., 5') from the DNA coding for this specific mRNA and that contains the promoter of this DNA.

[0040] Examples of a first promoter of this invention are the *Nicotiana tabacum* promoters of the STMG-type genes, described in Example 2, which are style and/or stigma specific promoters. Other style-stigma specific first promoters from other plant species can be isolated from their genomes, using the STMG-type genes as a probe as in step 4, above. Under hybridizing conditions, such a probe will hybridize to DNA coding for a style-stigma specific mRNA in a mixture of DNA sequences from the genome of the other plant species (Maniatis et al (1982) Molecular Cloning. A Laboratory Manual. Ed. Cold Spring Harbor Laboratory). Thereafter, as in step 5 above, another style-stigma specific first promoter can be identified. Other style-specific promoters can be isolated from self-incompatibility genes, such as an S-gen, for example as isolated from *Nicotiana glauca* (McClure et al (1989) Nature 342, 955-957). Other female organ-

specific promoters can be identified using other female organ-specific cDNAs, such as cDNA clone pMON9608 (Gasser et al (1989) *The Plant Cell* **1**, 15) that hybridizes exclusively with a gene expressed only in the ovules of tomato plants.

[0041] Other examples of such a first promoter are: the promoter of the KT13 gene (Jofuku and Goldberg (1989) *The Plant Cell* **1**, 1079-1093) which is an embryo axis-specific promoter; and the seed-specific promoters derived from genes encoding seed-specific storage proteins, such as the PAT2S promoters, for example PAT2S1, PAT2S2, PAT2S3 and PAT2S4 which are promoters of the four 2S albumin genes ("AT2S genes") of *Arabidopsis thaliana* (Krebbbers et al (1988) *Plant Physiol.* **87**, 859-866).

[0042] If more than one female-sterility DNA is present in the foreign DNA sequence of this invention, all the female-sterility DNAs can be under the control of a single first promoter, but preferably, each female-sterility DNA is under the control of its own separate first promoter. Where a plurality of female-sterility DNAs are present in the foreign DNA sequence, each female-sterility DNA can encode the same or different first RNA, polypeptide or protein. For example, when the female-sterility DNA encodes an RNase such as RNase T1, it is preferred that at least 3, particularly 4 to 6, copies of the female-sterility DNA and its first promoter be provided in the foreign DNA sequence. In such a case it is also preferred that all the female-sterility DNAs and their first promoters be adjacent to one another in the foreign DNA sequence and in any vector used to transform plant cells with the foreign DNA sequence. If the plurality of female-sterility DNAs encode different products, such as gene 1 and gene 2 or such as TNV replicase and a RNase, DNase or protease, it may be preferred that the female-sterility DNAs not be adjacent to one another and perhaps not even be present in the same vector used to transform plant cells with the foreign DNA sequence or even not present in the same parent plant of the female-sterile plant of this invention.

[0043] The selection of the marker DNA of this invention also is not critical.

[0044] In this regard the marker DNA may be any DNA sequence encoding a second RNA, protein or polypeptide which confers on at least a specific plant tissue or specific plant cells, in which such DNA sequence is expressed, a distinctive trait compared to such a specific plant tissue or specific plant cells in which such DNA sequence is not expressed.

[0045] A suitable marker DNA can be selected and isolated in a well known manner, so that it encodes a second RNA, protein or polypeptide that allows plants or their tissue, seeds or even cells, expressing the marker DNA, to be easily distinguished and separated from plants or their tissue, seeds or even cells not expressing the second RNA, protein or polypeptide. Examples of marker DNAs encode proteins that can provide a distinguishable color to plant cells, such as the A1 gene encoding dihydroquercetin-4-reductase (Meyer et al (1987) *Nature* **330**, 677-678) and the glucuronidase gene (Jefferson et al (1988) *Proc. Natl. Acad. Sci. USA* ("PNAS") **85**, 8447), or that provide a specific morphological characteristic to the plant such as dwarf growth or a different shape of the leaves. Other examples of marker DNAs confer on plants: stress tolerance, such as is provided by the gene encoding superoxide dismutase as described in European patent application 88/402222.9; disease or pest resistance such as is provided by a gene encoding a *Bacillus thuringiensis* endotoxin conferring insect resistance as described in European patent application 86/300291.1 or a gene encoding a bacterial peptide that confers a bacterial resistance as described in European patent application 88/401673.4.

[0046] Preferred marker DNAs encode second proteins or polypeptides inhibiting or neutralizing the action of herbicides such as: the *sfr* gene and the *sfrv* gene encoding enzymes conferring resistance to glutamine synthetase inhibitors such as bialaphos and phosphinotricine as described in European patent application 87/400,544.0; and genes encoding modified target enzymes for certain herbicides that have a lower affinity for the herbicides than naturally produced endogenous enzymes, such as a modified glutamine synthetase as a target for phosphinotricine as described in European patent publication 0,240,792 and a modified 5-enolpyruvylshikimate-3 phosphate synthase as a target for glyphosate as described in European patent publication 0,218,571. Other examples are: marker DNAs encoding proteins which neutralize the action of the herbicide bromoxynil (Stalker et al (1988) in: *Genetic Improvements of Agriculturally Important Crops*, Ed: R.T. Fraley, N.M. Frey and J. Schell, Cold Spring Harbor Laboratories); the herbicide sulfonylurea (Lee et al (1988) *EMBO J.* **7**, 1241-1248); and the herbicide 2,4 D (presented at the 2nd International Symposium of Plant Molecular Biology, Jerusalem, 13-18 November 1988).

[0047] The second promoter of this invention, which controls the marker DNA, can also be selected and isolated in a well known manner so that the marker DNA is expressed either selectively in one or more specific tissues or specific cells or constitutively in the entire plant, as desired depending on the nature of the second RNA, protein or polypeptide encoded by the marker DNA. For example, if the marker DNA encodes an herbicide resistance, it may be useful to have the marker DNA expressed in all cells of the plant, using a strong constitutive second promoter such as a 35S promoter (Odell et al (1985) *Nature* **313**, 810-812), a 35S'3 promoter (Hull and Howell (1987) *Virology* **86**, 482-493), the promoter of the nopaline synthetase gene ("PNOS") of the Ti-plasmid (Herrera-Estrella (1983) *Nature* **303**, 209-213) or the promoter of the octopine synthase gene ("POCS" [De Greve et al (1982) *J. Mol. Appl. Genet.* **1** (6), 499-511]). If the marker DNA encodes a protein conferring disease resistance, it may be useful to have the marker DNA selectively expressed in wound tissue by using, for example, a TR promoter such as the TR1' or TR2' promoter of the Ti-plasmid (Velten et al

(1984) EMBO J. 3, 2723-2730). If the marker DNA encodes a herbicide resistance, it also may be useful to have the marker DNA selectively expressed in green tissue by using, for example, the promoter of the gene encoding the small subunit of Rubisco (European patent application 87/400,544.0). If the marker DNA encodes a pigment, it also may be useful to have the marker DNA expressed in specific cells such as petal cells, leaf cells or seed cells, preferably in the outside layer of the seed coat.

[0048] One can identify and isolate in a well known manner a tissue-specific second promoter for a plant to be rendered female-sterile and easily distinguishable from non-transformed plants by:

1. searching for an mRNA which is only present in the plant during the development of a certain tissue, such as its petals, leaves or seeds;
2. isolating this tissue-specific mRNA;
3. preparing a cDNA from this tissue-specific mRNA;
4. using this cDNA as a probe to identify the regions in the plant genome which contain DNA coding for the tissue-specific mRNA; and then
5. identifying the portion of the plant genome that is upstream from the DNA coding for the tissue-specific mRNA and that contains the promoter for said DNA.

[0049] If more than one marker DNA is present in the foreign DNA sequence of this invention, all the marker DNAs can be under the control of a single second promoter, but preferably, each marker DNA is under the control of its own separate second promoter. More preferably, each marker DNA is under the control of its own second promoter and encodes a different second RNA, protein or polypeptide, providing different distinguishable characteristics to a transformed plant. In some cases, it may be preferred that the marker DNA(s) and second promoter(s) are adjacent to each other and to the one or more female-sterility DNAs contained in the foreign DNA sequence of this invention and in any vector used to transform plant cells with the foreign DNA sequence. In other cases, it may be preferred that the marker DNAs are not adjacent to each other and/or to the female-sterility DNAs.

[0050] It is generally preferred that the first RNA, protein or polypeptide, encoded by the female-sterility DNA, interferes significantly with the metabolism, functioning and/or development of the cells of the flowers and/or seeds and/or embryos by acting in the cytoplasm or the nucleus of these cells. However, when it is desired to have the first protein or polypeptide and/or the second protein or polypeptide transported from the cytoplasm into chloroplasts or mitochondria of the cells of transformed plants, the foreign DNA sequence can further include a first additional foreign DNA encoding a transit peptide. The first additional DNA is located between the female-sterility DNA and the first promoter if the first protein or polypeptide is to be so-transported and is between the marker DNA and the second promoter if the second protein or polypeptide is to be so-transported. By "transit peptide" is meant a polypeptide fragment which is normally associated with a chloroplast or mitochondrial protein or subunit of the protein that is produced in a cell as a precursor protein encoded by the nuclear DNA of the cell. The transit peptide is responsible for the translocation process of the nuclear-encoded chloroplast or mitochondrial protein or subunit into the chloroplast or the mitochondria, and during such a process, the transit peptide is separated or proteolytically removed from the chloroplast or mitochondrial protein or subunit. One or more of such first additional DNAs can be provided in the foreign DNA sequence of this invention for transporting one or more first or second proteins or polypeptides as generally described in European patent applications 85/402,596.2 and 88/402,222.9 and in: Van den Broeck et al (1985) Nature 313, 358-363; Schatz (1987) Eur. J. of Biochem. 165, 1-6; and Boutry et al (1987) Nature 328, 340-342. An example of a suitable transit peptide for transport into chloroplasts is the transit peptide of the small subunit of the enzyme RUBP carboxylase (European patent application 85/402,596.2), and an example of a transit peptide for transport into mitochondria is the transit peptide of the enzyme Mn-superoxide dismutase (see example 10 herein and European patent application 89/401,194.9).

[0051] It is also generally preferred that the first RNA, protein or polypeptide, encoded by the female-sterility DNA, act intracellularly so as to interfere with cell metabolism, functioning and/or development in the plant. However when it is desired to have the first protein or polypeptide and/or the second protein or polypeptide secreted out of the intercellular areas of the plant cells, in which they are expressed, or out of the tissue, in which they are expressed, the foreign DNA sequence can further include a second additional foreign DNA encoding a secretory signal peptide. The second additional foreign DNA is located between the female-sterility DNA and the first promoter if the first protein or polypeptide is to be secreted and between the marker DNA and the second promoter if the second protein or polypeptide is to be secreted. By "secretory signal peptide" is meant a natural polypeptide fragment which is, particularly in eukaryotic cells, associated during translocation with proteins that are normally secreted from cells or an artificial polypeptide fragment which, when associated during translocation with a protein or polypeptide, provokes its secretion from cells. Examples of suitable secretory signal peptides are set forth in: Von Heijne (1986), NAR 14 (11), 4683-4690; Denecke et al (1990), The Plant Cell 2, 51-59.

[0052] In the foreign DNA sequence of this invention, 3' transcription termination and polyadenylation signals can be selected in a conventional manner from among those which are capable of providing correct transcription termina-

tion and polyadenylation of mRNA in plant cells. The transcription termination and polyadenylation signals can be the natural ones of the gene to be transcribed but can also be foreign or heterologous. Examples of heterologous transcription termination and polyadenylation signals are those of the octopine synthase gene (Gielen et al (1984) EMBO J. 3, 835-845) and the T-DNA gene 7 (Velten and Schell (1985) Nucleic Acids Research ("NAR") 13, 6981-6998).

5 [0053] Also in accordance with this invention, plant cell cultures, containing the foreign DNA sequence of this invention, can be used to regenerate homozygous dominant female-sterile male-fertile plants by performing the necessary transformations on diploid (Chuong and B versdorf (1985) Plant Sci. 39, 219-226) or on haploid cell cultures and then (for haploid cell cultures) doubling the number of chromosomes by well known techniques (e.g., with colchicine). See: Plant Tissue and Cell Culture, Plant Biology 3, A.R. Liss, Inc. N.Y. (1987). Thereby, the foreign DNA sequence will be
10 in homozygous form in the nuclear genome of each of the so-transformed plant cells. This is preferred for plant cell cultures containing a female-sterility DNA under the control of a first promoter which directs gene expression at a given stage of development of the female gametes, such as ovums, especially after meiosis, or in cells derived from the female gametes, such as seed or embryo cells, so that the female-sterility DNA is present and can be expressed in all female gametes or plant cells derived therefrom.

15 [0054] Also in accordance with this invention, processes are provided for producing hybrid seeds which can be grown into hybrid plants. These processes involve crossing in an otherwise conventional manner: a) a nuclear female-sterile male-fertile plant of this invention, in which the first RNA, protein or polypeptide is expressed selectively in flowers, preferably in at least one female organ thereof, or in embryos; with b) a male-sterile female-fertile plant. Suitable male-sterile female-fertile plants are described in European Patent Application 89/401194.9 as having a nuclear
20 genome, in which are stably integrated:

- (a) a male-sterility DNA encoding a RNA, protein or polypeptide which, when produced or overproduced in a stamen cell of a plant, significantly disturbs adversely the metabolism, functioning and/or development of the stamen cell; and
- 25 (b) a promoter capable of directing expression of the male-sterility DNA selectively in stamen cells of the plant, preferably in anther, pollen and/or filament cells, particularly in tapetum and/or anther epidermal cells; the male-sterility DNA being in the same transcriptional unit as, and under the control of, this promoter;

and which optionally has in the same genetic locus:

- 30 (c) another marker DNA encoding a RNA, protein or polypeptide which, when present at least in a specific tissue or in at least specific cells of the plant, renders the plant easily separable from other plants which do not contain this RNA, protein or polypeptide at least in the specific tissue or specific cells; and
- (d) another promoter capable of directing expression of the other marker DNA at least in the specific tissue or specific cells and being in the same transcriptional unit as, and controlling, the other marker DNA.

The female-sterile plants and male-sterile plants are planted at random, near to each other to increase the chances of cross-pollination, without the need for precise planting patterns. The harvested seed, which is capable of germinating, will be the result of the fertilization of the male-sterile plants by the female-sterile plants and will be 100% hybrid. When
40 the foreign DNA sequences responsible for the female-sterility and male-sterility characteristics are present in heterozygous form in the nuclear genomes of the respective parent plants, plants grown from such hybrid seed will be: 25 % fertile, 25 % female-sterile, 25 % male-sterile and 25 % sterile. When the foreign DNA sequence encoding female-sterility is present in the nuclear genome of the male-fertile parent plant in homozygous form--which is preferred when the first promoter is an ovum-, seed- or embryo-specific promoter--all the plants grown from such hybrid seed will be
45 female-sterile.

[0055] Further in accordance with this invention, processes are provided for producing fruit without seeds by crossing in an otherwise conventional manner: a) a nuclear female-sterile male-fertile plant of this invention, in which the first RNA, protein or polypeptide is expressed selectively in seeds and in which the foreign DNA sequence, encoding the first RNA, protein or polypeptide, is preferably in homozygous form in the nuclear genome of the plant; with b) a male-sterile female-fertile plant.
50

[0056] Plants, transformed with the female-sterility DNA and in some cases preferably also with the marker DNA encoding an herbicide resistance, stably integrated in the plants' nuclear genomes and transmissible throughout generations as dominant alleles in accordance with this invention, are alternatives to, and can provide advantages over, presently used cytoplasmic and nuclear male-sterility systems for breeding and producing hybrid crops. In this regard,
55 female-sterile male-fertile plants can provide: 1) inhibited seed formation in crops, 2) hybrid seeds for crops which do not easily cross-pollinate, and 3) easier breeding of plant lines as discussed below.

1. Inhibition of seed formation

[0057] There exist a wide variety of crops cultivated by man in which the seed is an undesirable by-product.

5 a) When the economic product of a plant consists of its vegetative part. By inhibiting seed production, the plant's energy can be focused on vegetative biomass production. Examples are perennial plants (e.g. forage grasses, forage legumes and rubber trees), some annual plants (e.g., sugar cane and potato), and especially all crops that would normally flower and set seed before the economic product is harvested. Other examples are plants, obtainable through genetic engineering, which produce, within their vegetative tissues, proteins, polypeptides or other

10 metabolites for pharmaceutical or industrial purposes.
b) When the economic product of a plant is its fruit and it is desirable that the fruit be seedless, either because of consumer preferences (e.g., in tomato, melon and citrus fruit) or because seed formation uses up biomass that could otherwise be stored in the fruit (e.g., for providing high solids in tomatoes to be processed). Since such crops require fruit formation, the seedless condition, which usually induces fruit abortion, has to be compensated for by

15 the possibility of obtaining parthenocarpic fruit set. Natural parthenocarpic fruit inducing genes exist in some crops, such as tomato and melon.
c) When the plant is not grown for its seeds, but remaining propagules after harvesting may give rise to seed formation. The regrowth from these seeds can cause a considerable weed problem in the next culture. This "weed" problem is particularly well known with sugar beet.

20 d) When the plant is grown for its flowers (e.g., cut flowers, pot plants or garden ornamentals). For these species, it is often desirable to avoid seed set. In the case of cut flowers or pot plants, fertilization of the flowers often induces an accelerated senescence of the petals. In case of garden ornamentals, the formation of fruits and seeds reduces the time span and the intensity of flowering.

25 2. Hybrid seed production

[0058] Engineered female-sterility is useful as a seed production tool in combination with natural cytoplasmic or nuclear male-sterility systems or engineered nuclear male-sterility systems for the production of commercial hybrid seeds in crops where the seed is not the economic harvest and which do not easily cross-pollinate (e.g., for forage

A. Development of the female parent line A

[0059]

40 Aa) Transform line A with a foreign DNA including a male-sterility DNA under the control of a stamen specific promoter and adjacent thereto a marker DNA encoding herbicide resistance, according to European patent application 89/401,194.9, giving $A^{MSH/msh}$

Ab) Maintain line $A^{MSH/msh}$ through crossing with line $A^{msh/msh}$. This gives:

45 50% $A^{MSH/msh}$ (male-sterile, herbicide resistant) and
50% $A^{msh/msh}$ (fertile, herbicide sensitive).

B. Development of the male parent line B

[0060]

50 Ba) Transform line B with the chimaeric DNA sequence of this invention including a female-sterility DNA under the control of a first promoter which directs gene expression selectively in cells of a female organ of the plant and adjacent thereto a marker DNA encoding herbicide resistance, giving $B^{FSH/fsh}$.

Bb) Maintain line $B^{FSH/fsh}$ through crossing with $B^{fsh/fsh}$, yielding:

55 50% $B^{FSH/fsh}$ (female-sterile, herbicide resistant) and

50% B^{fst/fsh} (fertile, herbicide sensitiv).

C. Producing the hybrid seed crop

5 [0061]

Ca) Planting seeds obtained in Ab) and Bb) at random.

Cb) Eliminating through spraying with the herbicide the undesirable genotype before cross- and self-pollination could occur.

10 Cc) Cross-pollination occurring:

$A^{MSH/msh} \times B^{FSH/fsh}$

giving 100% hybrid seeds with the following genotype:

15

25% AB^{MSH/msh; FSH/fsh}

25% AB^{MSH/msh; fsh/fsh}

25% AB^{msh/msh; FSH/fsh}

25% AB^{msh/msh; fsh/fsh}

20

[0062] This represents the commercially sold seed.

3. Easier Breeding

25 a) without a marker DNA

[0063] The ability to obtain microspore-derived double haploids of most major crops allows the production of homozygous nuclear female-sterile lines in a more or less straightforward way (Chuong and Beversdorf (1985) Plant Sci. 39, 219-226). This makes it unnecessary in many cases to have a marker DNA within the same genetic locus of the nuclear genome of the cells of the plant as the female-sterility DNA. This is especially so if the homozygous female-sterile plant can be vegetatively multiplied (e.g., many vegetables).

30

b) with a marker DNA

35 [0064] In case the female-sterility DNA is in the same genetic locus of the nuclear genome of the transformed plant as a marker DNA (e.g., encoding herbicide resistance), homozygous female-sterile plants are technically superior to many other lines as tester parents in line evaluation programs. This is especially the case for crops where the seed is not the economic harvest and which can easily cross-pollinate. Indeed, these female-sterile plants allow the testing of many female- and male-fertile lines in close proximity to one another while making it easy to eliminate any self-pollinated seed from the different plant lines, being tested, from seeds resulting from crosses between the different lines.

40

[0065] The following Examples illustrate the invention. The figures referred to in the Examples are as follows:

Fig. 1A shows the cDNA sequence of the STMG07 gene of Example 1.

Fig. 1B shows the cDNA sequence of the STMG08 gene of Example 1.

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Fig. 2A shows the cDNA sequence of the STMG4B12 gene of Example 1.

Fig. 2B shows the cDNA sequence of the STMG3C9 gene of Example 1.

Fig. 3 shows a map of the vector pMG100 of Example 4.

Fig. 4 shows a map of the vector pMG101 of Example 6.

Fig. 5 shows a map of the vector pMG102 of Example 8.

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Fig. 6 shows a map of the vector pMG103 of Example 8.

Fig. 7 shows a map of the vector pMG104 of Example 10.

Fig. 8 shows a map of the vector pMG105 of Example 10.

[0066] Unless otherwise stated in the Examples, all procedures for making and manipulating recombinant DNA were carried out by the standardized procedures described in Maniatis et al, Molecular Cloning - A Laboratory Manual, Cold Spring Harbor Laboratory (1982). The following vector, used in the Examples, has been deposited in the Deutsche Sammlung Für Mikroorganismen und Zellkulturen ("DSMZ"), Mascheroder Weg 1B, D-3300 Braunschweig, Federal Republic of Germany under the provisions of the Budapest Treaty:

55

Vector	DSM Accession No.	Date
pGSC1700	4469	21 Mar. 1988

Example 1 - Isolation of style-stigma specific cDNAs from *Nicotiana tabacum* "Petit Havana" SR1.

[0067] Using well known procedures (Maniatis et al, 1982), total mRNA was isolated from the following different tobacco tissues: style-stigma tissues from flowers in stage 3 to 7 (according to Goldberg (1988) Science 240, 1460-1467); so-called "young stage" style-stigma tissues from a flower which did not develop pollen grains in stage 8 to 11 (according to Goldberg, 1988); so-called "old stage" ovary tissue from flowers in young stage; ovary tissue from flowers in old stage; and stem, root and leaf tissue from *in vitro* cultivated seedlings. cDNAs were synthesized from young and old style-stigma tissues using the Amersham (Amersham International PLC, Buckinghamshire, England) kit, cDNA Synthesis System Plus-RPN 1256 Y/Z, according to the directions set forth in the kit for its use. The cDNAs were cloned in Lambda gt 10 vector using the Amersham kit, cDNA Cloning System-lambda gt 10 - RPN 1257, according to the directions set forth in the kit. From the cDNA library thus obtained, differential screening was performed with a cDNA probe from seedlings on the one hand and a cDNA probe from style-stigma tissues on the other hand. The selected clones were subcloned in pGEM1 (Promega, Madison, Wisconsin, USA). Probes of each of these subclones were prepared and first checked for their specificity in Northern blots with 10 ug of total mRNA from different tobacco tissues (roots, stems, leaves, sepals, petals, anthers, young stage style-stigma, old stage style-stigma, and old stage ovaries). The subclones, that specifically hybridized in these Northern blots with style-stigma mRNA, were again hybridized in Northern blots with 2 ug poly A⁺ mRNA isolated from the above-mentioned tissues, including young ovaries, seeds and virus-infected leaves. The clones, called "pMG07" and "pMG08" and containing an insert of 0.963 kb and 0.472 kb, respectively, proved to be style-stigma specific cDNA sequences. These clones were sequenced, and their cDNA sequences are shown in Fig. 1A and Fig. 1B, respectively. The DNA sequence of pMG07 reveals the presence of one open reading frame ("ORF") over a sequence of 800 nucleotides. The sequence of pMG08 reveals an ORF over the total sequence.

[0068] From Prof. Goldberg of the University of California, Los Angeles (UCLA) were obtained: two *Nicotiana tabacum* style-stigma specific cDNAs (4B12 and 3C9) cloned as a *Pst*I - *Sma*I fragment in pGEM 3zf (-) (Promega, Madison, Wisconsin, USA). These clones contained inserts of 0.748 kb and 1.046 kb, respectively. Probes of these two clones were hybridized in Northern blots with 10 ug of total mRNA from different tobacco tissues (roots, stems, leaves, sepals, petals, anthers, young stage style-stigma, old stage style-stigma, and old stage ovaries) in order to check their specificity. These Northern blots confirmed the specificity of the clones and revealed that the transcript of 4B12 is 0.8 kb and that of 3C9 is 1.2 kb. The two clones were subcloned in pGEM1 (Promega), which subclones were called "pMG4B12" for the 4B12 clone and "pMG3C9" for the 3C9 clone. The subclones were again checked for their specificity in Northern blots with 2 ug poly A⁺ mRNA isolated from the above-mentioned tissues, including young ovaries, seeds and virus-infected leaves. The clones pMG4B12 and pMG3C9, containing inserts of 0.748 kb and 1.046 kb, respectively, proved to be style-stigma specific sequences. These clones were sequenced and their cDNA sequences are shown in Fig. 2A and Fig. 2B, respectively.

Example 2 - Isolation of the style-stigma specific genes ("STMG-type genes") corresponding to the style-stigma cDNA clones

[0069] Using known procedures (Maniatis et al, 1982), a probe from each of the cDNA clones of Example 1 of style-stigma specific sequences was used to isolate the corresponding genomic gene sequence which is specifically expressed in style-stigma tissues of the female organ of tobacco. According to protocols provided by Promega, tobacco genomic DNA was partially digested with *Sau*3A, and the restriction fragments were cloned into the lambda phage vector GEM.12 (Promega), digested with *Xho*I to produce genomic clones called "lambda STG07", "lambda STG08", "lambda STG4B12" and "lambda STG3C9". Subsequently, these genomic clones were subcloned in pGEM1 (Promega) according to the procedure of Promega. The subclones were again analyzed by Southern blot, using the

respective cDNA clones as probes in order to identify the clones which contained the style-stigma specific DNA sequences. These subclones, called respectively "pSTG07", "pSTG08", "pSTG4B12" and "pSTG3C9" were sequenced (Maxam and Gilbert (1977) PNAS 74, 560). The orientation of these clones was determined by Northern blot analysis with riboprobes of both senses. Comparison of each cDNA sequence with its respective genomic clone sequence led to the identification of the region of homology. At the 5' end of each region, the ATG codon and the consensus sequence TATA were determined. That the "TATA" box is part of the promoter of the gene is confirmed by primer extension (Mcknight et al (1987) Cell 25, 385). The style-stigma specific genes, isolated using the style-stigma cDNA as probe, are called in general "STMG-type" genes. The style-stigma specific gene of pSTG07 is called "STMG07", that of pSTG08 is called "STMG08", that of pSTG4B12 is called "STMG4B12" and that of pSTG3C9 is called "STMG3C9".

Example 3 - Construction of promoter cassettes ("PSTMG") derived from the respective STMG-type genes

[0070] To construct chimaeric DNA sequences containing 5' regulator sequences, including the promoter of an STMG-type gene, in the same transcriptional unit as, and controlling, a first heterologous female-sterility DNA, cassettes are constructed by subcloning a DNA fragment including a promoter into the polylinker of the pMAC5-8 vector system (European patent application 87/402348.4). This produces respective vectors which can be used to isolate single strand DNA for use in site directed mutagenesis.

[0071] Using site directed mutagenesis (European patent application 87/402348.4), the sequence surrounding the ATG initiation codon of each of the genes is modified in such a way that the mutation creates a given sequence which is a unique recognition site for a given restriction enzyme. The resulting plasmids each contain the newly created restriction site. The precise nucleotide sequence spanning the newly created restriction site is determined in order to confirm that it only differs from the 5' sequence of the corresponding STMG-type gene by the substitution, creating the new restriction site. The newly created promoter cassettes, each comprising a promoter, a 5' untranslated end of an STMG-type gene to its ATG initiation codon, and a new restriction site, are generally called "PSTMGs". The PSTMG containing the promoter and 5' end of STMG07 is called "PSTMG07", that of STMG08 is called "PSTMG08", that of STMG4B12 is called "PSTMG4B12" and that of STMG3C9 is called "PSTMG3C9".

Example 4 - Construction of chimaeric DNA sequences of a PSTMG and a RNase T1 gene

[0072] Plasmids named "pMG100", shown in Fig. 3, are constructed, each by assembling the following well known DNA fragments with a different one of the PSTMGs of Example 3:

1. a vector fragment, including T-DNA border sequences, from pGSC1700 in which the β -lactamase gene encoding ampicillin has been inactivated by insertion into the SacI site; located between the border sequences are the following DNA fragments 2-4;
2. a chimaeric sequence containing an Arabidopsis Rubisco SSU promoter ("PSSU" or "PSSUARA"), a herbicide resistance gene sfr (European patent application 87/400,544.0) and the 3' end (i.e., transcription termination) signals of a T-DNA gene 7 (Velten and Schell (1985) NAR 13, 6981);
3. a chimaeric sequence containing the EcoRI/SacI fragment from pGSFR401 which contains a nopaline-synthase promoter ("PNOS"), a neo gene encoding kanamycin resistance and the 3' end signals of an octopine synthase ("OCS") gene (European patent application 87/400,544.0, wherein pGSFR401 is called "pGSR4"); and
4. a chimaeric sequence, containing one of the PSTMG promoter cassettes from Example 3, fused in frame with a synthetic gene encoding RNase T1 from A. oryzae, (Quaas et al, "Biophosphates and their Analogues-Synthesis, Structure, Metabolism and Activity" (1987) Elsevier Science Publisher B.V., Amsterdam; Quaas et al (1988) Eur. J. Biochem. 173, 617-622) and the 3' end signals of a nopaline synthase ("NOS") gene (An et al (1985) EMBO J. 4 (2), 277).

[0073] Each pMG100 is a binary type T-DNA vector containing, within the T-DNA border sequences, three chimaeric sequences: PSSU-sfr and PNOS-neo which are marker DNAs with respectively PSSU and PNOS as second promoters; and PSTMG-RNase T1 gene which is a female-sterility DNA under the control of a PSTMG as a first promoter. Expression of the female-sterility DNA under the control of the PSTMG promoter will produce RNase T1 selectively in style and/or stigma cells. This will be lethal for the style and/or stigma cells since the RNase T1 will degrade the RNA molecules which are indispensable for these cells' metabolism.

Example 5 - Introduction of each chimaeric DNA sequence of Example 4 into tobacco and alfalfa

[0074] A recombinant Agrobacterium strain is constructed by mobilization of each pMG100 (from Example 4) from E. coli into Agrobacterium tumefaciens C58C1 Rif^R containing pMP90 (Koncz and Schell (1986) Mol. Gen. Genetics

204, 382-396).

[0075] The resulting *Agrobacterium* strain, harboring pMP90 and pMG100, is used for the transformation of tobacco leaf discs (*N. tabacum* Petite Havane SR1), using standard procedures as described, for example, in European patent application 87/400,544.0, and of alfalfa according to the procedure described in D'Halluin et al (1990) Crop Science 30:866-871. Carbenicillin is used to kill the *Agrobacterium* strains after co-cultivation. Transformed calli are selected on a substrate containing 5 mg/l phosphinotricin and 100 ug/ml kanamycin, and resistant calli are regenerated into plants. After induction of shoots and roots, which proves normal growth of the plants despite the presence of the RNase T1 gene, the transformants are transferred to the greenhouse and are grown until they flower. The flowers are examined and show no normal style-stigma formation. After pollination, no viable seeds are formed. The transformed plants are female-sterile.

Example 6 - Construction of chimaeric DNA sequences of a PSTMG and a Barnase gene

[0076] Plasmids named "pmg101", shown in Fig.4, are constructed, each by assembling the following well known DNA fragments with a different one of the PSTMGs of Example 3:

1. a vector fragment, including T-DNA border sequences, derived from pGSC1700 as described in Example 4 and with the following DNA fragments 2-4 between its border sequences;
2. the chimaeric sequence (no. 2) of Example 4, containing the PSSU promoter, the herbicide-resistance gene *sfr* and the 3' end of T-DNA gene 7;
3. the chimaeric sequence (no. 3) of Example 4, containing the PNOS promoter, the *neo* gene encoding kanamycin resistance and the 3' end signals of the OCS gene; and
4. a chimaeric sequence, containing one of the PSTMGs from Example 3, fused in frame with the Barnase gene from *Bacillus amyloliquefaciens* (Hartley and Rogerson (1972) Preparative Biochemistry 2 (3), 243-250) and the 3' end of the NOS gene of Example 4.

[0077] Each pMG101 is a binary type T-DNA vector containing, within the T-DNA border sequences, three chimaeric sequences: PSSU-*sfr* and PNOS-*neo* which are markers DNAs with respectively PSSU and PNOS as second promoters; and PSTMG-Barnase gene which is a female-sterility DNA under the control of a PSTMG as a first promoter. Expression of the female-sterility DNA under the control of the PSTMG promoter will produce Barnase selectively in style and/or stigma cells. This will be lethal for the style and/or stigma cells since Barnase will degrade the RNA molecules and thereby interfere with the metabolism of these cells.

Example 7 - Introduction of each chimaeric DNA sequence of Example 6 into tobacco and alfalfa

[0078] As described in Example 5, a recombinant *Agrobacterium* strain is constructed by mobilizing each pMG101 (from Example 6) from *E. coli* into *Agrobacterium* C58C1 Rif^R containing pMP90 (Koncz and Schell (1986) Mol. Gen. Genetics 204, 383-396). The resulting strain, harboring pMP90 and pMG101, is used for tobacco leaf disc transformation and for alfalfa transformation. Transformed calli and shoots are selected using 5mg/l phosphinotricin and 100 ug/ml kanamycin. That the Barnase gene is not expressed in the transformed herbicide-resistant calli and shoots is shown by their growth.

[0079] The transformed shoots are rooted, transferred to soil in the greenhouse and grown until they flower. The flowers of both the tobacco and alfalfa are examined, and essentially the same phenotype is observed in the transformed plants as is observed in the transformed plants described in Example 5 (i.e., no normal style-stigma formation). The transformed plants are female-sterile.

Example 8 - Construction of chimaeric DNA sequences of a PSTMG and a gene encoding papain

[0080] Plasmids named "pmg102", shown in Fig 5, are constructed, each by assembling the following well known DNA fragments with a different one of the PSTMGs of Example 3:

1. a vector fragment, including T-DNA border sequences, derived from pGSC1700 as described in Example 4 and with the following DNA fragments 2-4 between its border sequences;
2. the chimaeric sequence (no. 2) of Example 4, containing the PSSU promoter, the herbicide resistance gene *sfr* and the 3' end of T-DNA gene 7.
3. the chimaeric sequence (no. 3) of Example 4, containing the PNOS promoter, the *neo* gene and the 3' end of the OCS gene; and
4. a chimaeric sequence, containing one of the PSTMGs from Example 3, fused in frame with:

a) a papain gene from Carica papaya fruit, encoding the papain zymogen which is a plant endopeptidase (Cohen et al (1986) Gene 48, 219-227) capable of attacking peptide, as well as ester, bonds; the following modifications are made in the DNA sequence of the papain gene according to Cohen et al (1986), using site directed mutagenesis as described in Example 3:

- i. the nucleotide A, position-1 upstream of the first ATG codon, is mutated into nucleotide C in order to obtain a suitable NcoI cloning site; and
- ii. the GAA codons encoding glutamate at positions 47, 118 and 135 are mutated into CAA codons encoding glutamine; and

b) the 3' end of the NOS gene of Example 4.

[0081] Each pMG102 is a binary type T-DNA vector containing, within the T-DNA border sequences, three chimaeric sequences: PSSU-sfr and PNOS-neo which are marker DNAs encoding dominant selectable markers for plant transformation under the control of respectively PSSU and PNOS as second promoters; and PSTMG-Papain zymogen gene which is a female-sterility DNA under the control of a PSTMG as a first promoter. Expression of the female-sterility DNA under the control of the PSTMG promoter will produce, selectively in style and/or stigma cells, an endopeptidase (the papain zymogen) that will cleave proteins in the style and/or stigma cells, thus leading to the death of these cells.

[0082] Plasmids named "pMG103", shown in Fig.6, are also constructed, each by assembling the following well known DNA fragments with a different one of the PSTMGs of Example 3:

1. a vector fragment, including T-DNA border sequences, derived from pGSC1700 as described in Example 4 and with the following DNA fragments 2-4 between its border sequences;
2. the chimaeric sequence (no. 2) of Example 4, containing the PSSU promoter, the herbicide resistance gene sfr and the 3' end of T-DNA gene 7;
3. the chimaeric sequence (no. 3) of Example 4, containing the PNOS promoter, the neo gene, and the 3' end of the OCS gene; and
4. a chimaeric sequence, containing one of the PSTMGs of Example 3, fused in frame with:

a) a papain gene from Carica papaya fruit, encoding the active protein of the papain zymogen; the following modifications are made in the DNA sequence of the papain gene according to Cohen et al (1986), using site directed mutagenesis as described in Example 3:

- i. the AAT codon encoding Asn, upstream of the first Ile residue of the active protein, is mutated into a GAT codon, which provides a suitable EcoRV cloning site (GAT ATC). The EcoRV engineered site is fused directly to the PSTMG in order to obtain a direct in frame fusion of the promoter with the sequence encoding the active protein of the papain zymogen; and
- ii. the GAA codons encoding glutamate at positions 47, 118 and 135 are mutated into CAA codons encoding glutamine; and

b) the 3' end of the NOS gene of Example 4.

[0083] Each pMG103, like each pMG102, is a binary type T-DNA vector containing, within the T-DNA border sequences, three chimaeric genes: PSSU-sfr and PNOS-neo encoding dominant selectable markers for plant transformation; and PSTMG-Papain active protein gene which is a female-sterility DNA that is under the control of a PSTMG as a first promoter and that encodes an endopeptidase that will cleave proteins in style and/or stigma cells, thus leading selectively to the death of these cells.

Example 9 - Introduction of each chimaeric DNA sequence of Example 8 into tobacco and alfalfa

[0084] As described in Example 5, each pMG102 and pMG103 (from Example 8) is mobilized from E. coli into separate Agrobacteria C58C1 Rif^R carrying pMP90. The resulting strains, harboring pMP90 with pMG102 and pMP90 with pMG103, are used to transform tobacco and alfalfa following the procedures of Example 5. That the papain genes are not expressed in transformed herbicide- and kanamycin-resistant calli, shoots and roots is shown by their growth.

[0085] The transformed plants are transferred into the greenhouse and grown in soil until they flower. The flowers of both the tobacco and alfalfa are examined, and essentially the same phenotypes are observed in the transformed plants as are observed in the transformed plants described in Example 5 (i.e., no normal style-stigma formation). The transformed plants are female-sterile.

Exempl 10 - Construction of chima ric DNA sequences f a PSTMG and a g ne enc ding EcoRI

[0086] Plasmids named "pMG104", shown in Fig. 7, are constructed, each by assembling the following well known DNA fragments with a different one of the PSTMGs of Example 3:

1. a vector fragment, including T-DNA border sequences, derived from pGSC1701A2 (European patent application 87/115985.1); located between the border sequences are the following DNA fragments 2-5;
2. the chimaeric sequence (no. 2) of Example 4, containing the PSSU promoter, the herbicide-resistance gene sfr and the 3' end of T-DNA gene 7;
3. the chimaeric sequence (no. 3) of Example 4, containing the PNOS promoter, the neo gene and the 3' end of the OCS gene;
4. a chimaeric sequence, containing one of the PSTMGs of Example 3, fused in frame with:

a) a gene encoding the EcoRI restriction endonuclease from an E. coli (Green et al (1981) J. Biol. Chem. 256, 2143-2153; Botterman and Zabeau (1985) Gene 37, 229-239) and capable of recognizing and cleaving the target sequence GAATTC on a double stranded DNA; the following modifications are made in the DNA sequence of the gene according to Green et al (1981) using site directed mutagenesis as described in Example 3:

- i. the nucleotides of the ATG initiation codon are replaced by ATGCA, creating a NsiI site at the initiation codon and yielding the following nucleotide sequences: ATGCA, TCT, AAT...; and
- ii. the HindII-HindIII fragment of the EcoRI gene cloned in pEcoR12 (Botterman and Zabeau, 1985) is cloned into the pMAC5-8 site directed mutagenesis vector; and

b) the 3' end of the NOS gene of Example 4; and

5. a gene encoding an EcoRI methylase, under the control of its natural promoter (Botterman and Zabeau, 1985), which is capable of inhibiting the activity of EcoRI in E. coli or Agrobacterium, in order to overcome potential leaky expression of the EcoRI gene in microorganisms.

[0087] Each pMG104 is a binary type T-DNA vector containing, within the T-DNA border sequences, three chimaeric sequences: PSSU-sfr and PNOS-neo which are marker DNAs under the control of respectively PSSU and PNOS as second promoters; and PSTMG-EcoRI endonuclease gene which is a female-sterility DNA under the control of a PSTMG as a first promoter. Expression of the female-sterility DNA under the control of the PSTMG promoter selectively in style and/or stigma cells will produce the EcoRI restriction endonuclease which will cleave double stranded DNA at GAATTC sites (see for review of type II restriction modification systems: Wilson (1988) TIG 4 (11), 314-318) in the style and/or stigma cells, thus leading to the death of these cells.

[0088] Plasmids named "pMG105" are also constructed, each by assembling the following well known DNA fragments with a different one of the PSTMGs of Example 3:

1. a vector fragment, including T-DNA border sequences, derived from pGSC1701A2; located between the border sequences are the following DNA fragments 2-5;
2. the chimaeric sequence (no. 2) of Example 4, containing the PSSU promoter, the herbicide-resistance gene sfr and the 3' end of T-DNA gene 7;
3. the chimaeric sequence (no. 3) of Example 4, containing the PNOS promoter, the neo gene and the neo 3' end of the OCS gene;
4. a chimaeric sequence, containing one of the PSTMGs of Example 3, fused in frame with:

a) a gene fragment encoding the transit peptide of the Mn-superoxide dismutase ("Mn-SOD") which is a NcoI-PstI fragment of a HpaI-HindIII fragment from pSOD1 (Bowler et al (1989) Embo J. 8, 31-38); the following modifications are made in the DNA sequence of the gene fragment according to Bowler et al (1989) using site directed mutagenesis as described in Example 3:

- i. the AA nucleotides located upstream at position -2 and -1 of the ATG initiation codon are changed to CC nucleotides creating a NcoI site at the initiation codon and yielding the following nucleotide sequences:

- CCATGGCACTAC

NcoI

5

10

ii. the T,TCG,CTC nucleotides located immediately downstream of the processing site of the transit peptide are changed to C,TGC,AGC, creating a PstI site behind the processing site and yielding the following nucleotide sequences:

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		L	Q	T	F	S	L
CTC, CGC, GGC,		TTG, CAG, ACC, TTT, TCG, CTC					
CTC, CGC, GGC,		TTG, CAG, ACC, <u>TTC, TGC, AGC</u>					
						<u>PstI</u>	

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in which the arrow indicates the processing site of the transit peptide sequence and the upper line the amino acid sequence corresponding with the Mn-SOD coding sequence; the NcoI-PstI fragment is also fused in frame with a gene encoding the EcoRI restriction endonuclease from E. coli (Greene et al (1981) J. Biol. Chem. 256, 2143-2153; Botterman and Zabeau (1985) Gene 37, 229-239) and capable of recognition and cleavage of the target sequence GAATTC on a double stranded DNA, as found in pMG104; and

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b) the 3' end of the NOS gene of Example 4; and

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5. a gene encoding the EcoRI methylase under the control of its natural promoter (Botterman and Zabeau, 1985), which is capable of inhibiting the activity of EcoRI in E. coli or Agrobacterium, in order to overcome potential leaky expression of the EcoRI gene in microorganisms, this gene being inserted into the vector fragment outside the border sequences.

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[0089] Each pMG105 is a binary type T-DNA vector containing, within the border sequences, three chimeric sequences: PSSU-sfr and PNOS-NPTII which are marker DNAs under the control of respectively PSSU and PNOS as second promoters; and PSTMG-transit peptide-EcoRI endonuclease gene which is a female-sterility DNA having the PSTMG as a first promoter and a transit peptide-encoding sequence between them. Expression of the female-sterility DNA under the control of the PSTMG promoter selectively in style and/or stigma cells will produce a restriction endonuclease which will be targeted into the mitochondria of the style and/or stigma cells and cleave the double stranded DNA at the GAATTC sites in such cells. This will lead to the death of these cells.

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Example 11 - Introduction of each chimaeric DNA sequence of Example 10 into tobacco and alfalfa

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[0090] As described in Example 5, each pMG104 and pMG105 (from Example 10) is mobilized from E. coli into separate Agrobacteria C58C1 Rif^R carrying pMP90. The resulting strains, harboring pMG104 with pMP90 and pMG105 with pMP90, are used to transform tobacco and alfalfa following the procedures described in Examples 5. That the EcoRI endonuclease genes are not expressed in transformed herbicide- and kanamycin-resistant calli, shoots and roots is shown by their growth.

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[0091] The transformed plants are transferred into the greenhouse and grown in soil until they flower. The flowers of both the tobacco and alfalfa are examined, and essentially the same phenotypes are observed for the transformed plants as are observed in the transformed plants described in Example 5 (i.e., no normal style-stigma formation). The transformed plants are female-sterile.

Claims

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1. A DNA which contains a first chimeric DNA which comprises:

(a) a female sterility DNA encoding a first RNA, protein or polypeptide, capable, when produced in cells of a female reproductive organ of said plant, of disturbing significantly the metabolism, functioning and/or develop-

ment of the female reproductive organ; and

(b) a first promoter capable of directing expression of this DNA selectively in cells of the female reproductive organs of said plant, said female sterility DNA being in the same transcriptional unit as, and under the control of said first promoter.

2. The DNA of claim 1 in which said first promoter is capable of directing expression of said female sterility DNA especially in style cells, stigma cells, ovary cells, ovule cells and/or septum cells.

3. The DNA of claim 1, wherein said first promoter is capable of directing expression of said female-sterility DNA especially in style and/or stigma cells.

4. The DNA of any one of claims 1 to 3 wherein said first promoter is selected from the group consisting of : PSTMG07 which is the promoter of an endogenous tobacco gene encoding an mRNA from which the cDNA of Fig. 1A can be prepared; PSTMG08 which is the promoter of an endogenous tobacco gene encoding an mRNA from which the cDNA of Fig. 1B can be prepared; PSTMG4B12 which is the promoter of an endogenous tobacco gene encoding an mRNA from which the cDNA of Fig. 2A can be prepared; PSTMG3C9 which is the promoter of an endogenous tobacco gene encoding an mRNA from which the cDNA of Fig. 2B can be prepared.

5. The DNA of any one of claims 1 to 4 wherein said first promoter is a promoter of a gene corresponding to cDNA clone pMON9608.

6. The DNA of any one of claims 1 to 5 wherein said female-sterility DNA encodes a ribonuclease, particularly RNase T1.

7. The DNA of any one of claims 1 to 5 wherein said female-sterility DNA encodes a Barnase

8. The DNA of any one of claims 1 to 5 wherein said female-sterility DNA encodes a DNase, especially an endonuclease, particularly EcoRI; a protease especially a papain, particularly papain Zymogen or papain active protein; a glucanase; a lipase, particularly phospholipase A₂; a lipid peroxidase; a cell wall inhibitor; a bacterial toxin; or a ribozyme; particularly a ribozyme against mRNA encoded by any of the STMG-type genes, or by a gene corresponding to cDNA clone pMON9608 ; or alternatively, wherein said female-sterility DNA: encodes a ribonuclease, particularly T₂ or Rh; encodes a glycoprotein encoded by the S1, S2, S3, S6 or S7 alleles, particularly of Nicotiana glauca; or is an anti-sense DNA, particularly an anti-sense DNA encoding an RNA which is complementary to the mRNA of an STMG-type gene.

9. The DNA of any one of claims 1 to 5, wherein said female-sterility DNA encodes an enzyme which catalyzes the synthesis of a phytohormone, particularly the enzyme encoded by gene 4 of Aprobacterium T-DNA, or an enzyme encoded by gene 1 and/or gene 2 of Aprobacterium T-DNA.

10. The DNA of any one of claims 1 to 5 wherein said female-sterility DNA encodes a viral dependant RNA polymerase, particularly a TNV replicase, under the control of said first promoter; and a negative strand of a gene encoding said first protein or polypeptide; said gene being in the same transcriptional unit as, and under the control of, a promoter, such as another first promoter, capable of directing expression of said negative strand in the cell; said gene also being fused at its 3' end to a viral subgenomic promoter, such as the TNV subgenomic promoter, which, is specifically recognized by said viral dependent RNA polymerase.

11. The DNA of any one of claims 1 to 10 wherein said DNA further comprises:

(e) a first DNA encoding a transit peptide capable of transporting said first protein or polypeptide into a chloroplast or mitochondria of said female reproductive organ cells: said first DNA being in the same transcriptional unit as said female-sterility DNA and said first promoter and between said female-sterility DNA and said first promoter.

12. The DNA of any one of claims 1 to 10 wherein said DNA further comprises:

(e) a third DNA encoding a secretory signal peptide capable of secreting said first protein or polypeptide outside the cell of a female reproductive organ of a plant; said third DNA being in the same transcriptional unit as said female-sterility DNA and said first promoter and located between said female-sterility DNA and said first

promoter.

13. The DNA of anyone of claims 1 to 12 which also comprises a second chimeric DNA, adjacent to said first chimeric DNA, said second chimeric DNA comprising:

(c) a marker DNA encoding a second RNA, protein or polypeptide which, when present at least in a specific tissue or in at least specific cells of a plant, renders said plant easily separable from other plants which do not contain said second RNA, protein or polypeptide in said specific tissue or specific cells; and
(d) a second promoter capable of directing expression of said marker DNA at least in said specific tissue or specific cells: said marker DNA being in the same transcriptional unit as, and under the control of, said second promoter.

14. The DNA of claims 13 wherein said marker DNA encodes a protein inhibiting or neutralizing the action of a herbicide.

15. The DNA of claim 14 in which said marker DNA is an herbicide resistance gene, particularly a gene conferring resistance to a glutamine synthetase inhibitor, such as phosphinothricin.

16. The DNA of claim 15 in which said marker DNA is a *sfr* or *sfrv* gene.

17. The DNA of claim 14 wherein said marker DNA is a gene encoding a modified target enzyme for an herbicide having lower affinity for the herbicide, particularly a modified 5-enolpyruvylshikimate-3 phosphate synthase as a target for glyphosate or a modified glutamine synthetase as a target for a glutamine synthetase inhibitor such as phosphinothricin.

18. The DNA of claim 13, wherein said marker DNA is a gene encoding a protein or a polypeptide conferring a color to at least said specific tissue or specific cells, particularly the gene *A1* or the *GUS* gene; a gene encoding a protein or a polypeptide conferring a stress tolerance to said plant, particularly the gene encoding Mn-superoxide dismutase; or a gene encoding a protein or a polypeptide conferring a disease or pest resistance, particularly a gene encoding a *Bacillus thuringiensis* endotoxin that confers insect resistance or a gene encoding a bactericidal peptide that confers a bacterial resistance.

19. The DNA of any one of claims 13 to 18 wherein said second promoter is: a constitutive promoter, particularly a *PNOS* promoter or a *POCS* promoter; a wound-inducible promoter, particularly a *TR1'* or *TR2'* promoter; a promoter which directs gene expression selectively in plant tissue having photosynthetic activity, or a promoter which directs gene expression selectively in leaf cells, petal cells or seed cells, particularly seed coat cells.

20. The DNA of claim 19 wherein said second promoter is a *35S* promoter or an *SSU* promoter.

21. The DNA of any one of claims 13 to 20 wherein said DNA further comprises:

(h) a second DNA encoding a transit peptide capable of transporting said second protein or polypeptide into a chloroplast or mitochondria of at least said specific tissue or specific cells; said second DNA being in the same transcriptional unit as said marker DNA and said second promoter and between said marker DNA and said second promoter.

22. The DNA of any one of claims 13 to 20 wherein said DNA further comprises:

(f) a fourth DNA encoding a secretory signal peptide capable of secreting said second protein or polypeptide outside at least specific tissue or specific cells; said fourth DNA being in the same transcriptional unit as said marker DNA and said second promoter and located between said marker DNA and said second promoter.

23. A DNA which comprises the T-DNA of pMG100 of Fig. 3, pMG101 of Fig. 4, pMG102 of Fig. 5, pMG103 of Fig. 6, pMG104 of Fig. 7, or pMG105 of Fig. 8.

24. The DNA of any one of claims 1 to 23 which is nuclear DNA of a cell of a plant or of a seed.

25. A vector which contains the DNA of any one of claims 1 to 23.

26. A cell of a plant which contains the DNA of any one of claims 1 to 24.
27. A cell of a plant which contains, stably integrated into its nuclear DNA, the DNA of any one of claims 1 to 23.
- 5 28. A cell of claim 26 or claim 27 which can be regenerated into a plant which is female-sterile.
29. A plant cell culture containing plant cells of any one of claims 26 to 28.
30. A plant which contains the DNA of any one of claims 1 to 24.
- 10 31. A plant which contains the DNA of any one of claims 1 to 24 in all of its cells.
32. The plant of claim 30 or 31 which is female-sterile.
- 15 33. The plant of any one of claims 30 to 32 which is a hybrid plant.
34. A female-sterile plant containing a foreign DNA incorporated in the nuclear genome of its cells, wherein said foreign DNA comprises :
- 20 (a) a female-sterility DNA encoding a first RNA, protein or polypeptide, capable when produced in cells of a female reproductive organ of said plant, of disturbing significantly the metabolism, functioning and/or development of the female reproductive organ; and
- (b) a first promoter capable of directing expression of this DNA selectively in cells of the female reproductive organs of said plant, said female sterility DNA being in the same transcriptional unit as, and under the control
- 25 of said first promoter,
- provided that, if said first promoter is a promoter capable of directing expression of said female-sterility DNA selectively in female gametes, or in cells derived from female gametes, the nuclear genome of said plant is homozygous.
- 30 35. The plant of anyone of claims 30 to 34 which is selected from the group consisting of alfalfa and tobacco.
36. A plant seed which contains the DNA of any one of claims 1 to 24.
37. A seedless fruit of a plant which contains the DNA of any one of claims 1 to 24.
- 35 38. A process for producing a female-sterile plant and reproductive material for said plant, which comprises :
- a) introducing the foreign DNA of any one of claims 1 to 23 into the nuclear genome of a plant cell to obtain a transformed plant cell;
- 40 b) regenerating said female-sterile plant from said transformed plant cell and; optionally,
- c) obtaining said reproductive material from said female-sterile plant which contain said foreign DNA.
39. The process of claim 20 in which said reproductive material is a seed, particularly a hybrid seed, and which further comprises :
- 45 a) providing a plant which contains the DNA of any one of claims 1 to 23 incorporated in the nuclear DNA of all of its cells and which is female-sterile;
- b) cross-pollinating i) said female-sterile plant, and ii) a female-fertile plant, and
- 50 c) recovering said seed from said female-fertile plant.
40. The process of claim 20 in which said reproductive material is a seed, particularly a hybrid seed, which comprises the steps of :
- 55 a) providing a plant which contains the DNA of any one of claims 13 to 23, including both said second promoter and said marker DNA, incorporated in the nuclear DNA of all of its cells, and which is female-sterile;
- b) cross-pollinating i) said female-sterile plant, and ii) a female-fertile plant without said marker DNA and/or said second promoter, and
- c) recovering the seed from said female-fertile plant,

wherein said process optionally comprises removing undesired female-fertile plants on the basis of the absence of expression of the marker DNA in said female-fertile plants.

41. A process for producing a seed which comprises the steps of:

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(a) providing a plant which contains the DNA of any one of claims 13 to 23 incorporated in the nuclear DNA of all of its cells, which is female-sterile, and in which said marker DNA is a herbicide resistance gene, particularly a gene conferring resistance to a glutamine synthetase inhibitor such as the *sfr* or *sfrv* gene, or a gene encoding a modified target enzyme for a herbicide having lower affinity for a herbicide, particularly a modified 5-enolpyruvylshikimate-3 phosphate synthase as a target for glyphosate or a modified glutamine synthetase as a target for a glutamine synthase inhibitor such as phosphinothricin,

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(b) cross-pollinating i) said female-sterile plant, and ii) a female-fertile plant without said marker DNA and/or said second promoter, and

(c) recovering the seed from said female-fertile plant,

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wherein said process comprises removing undesired female-fertile plants by application of said herbicide, particularly a herbicide comprising phosphinothricin.

42. A process according to claim 41 in which said seed is a hybrid seed.

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43. The process of anyone of claims 41 or 42, wherein said female-sterile plant contains in addition to said herbicide resistance gene at least another marker DNA stably integrated into the nuclear genome of its cells in the same genetic locus as said female-sterility DNA and said female-fertile plant does not contain said another marker DNA.

25 44. A process according to anyone of claims 39 to 43, in which the female-fertile plant is male-sterile.

45. A process according to claim 44, wherein said female-fertile male-sterile plant comprises a foreign DNA incorporated in the nuclear genome of its cells, wherein said foreign DNA comprises:

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(a) a male sterility DNA encoding a third RNA, protein or polypeptide, capable when produced in stamen cells of said plant, of disturbing significantly the metabolism, functioning and/or development of the stamen cells.

(b) a third promoter capable of directing expression of this DNA selectively in stamen cells of said plant, said male sterility DNA being in the same transcriptional unit as, and under the control of said third promoter;

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provided that, if said third promoter is a promoter capable of directing expression of said male-sterility DNA selectively in pollen cells, the nuclear genome of said transformed plant cell is homozygous.

46. A process according to claim 45, wherein said foreign DNA of said female-fertile male-sterile plant also comprises :

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(d) a second marker DNA encoding a fourth RNA, protein or polypeptide which, when present at least in a specific tissue or in at least specific cells of said plant, renders said plant easily separable from other plants which do not contain said fourth RNA, protein or polypeptide in said specific tissue or specific cells; and

(e) a fourth promoter capable of directing expression of said second marker DNA at least in said specific tissue or specific cells; said second marker DNA being in the same transcriptional unit as, and under the control of,

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said fourth promoter.

47. A process according to anyone of claims 45 or 46, wherein said foreign DNA further comprises :

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(f) a fifth DNA encoding a transit peptide capable of transporting said third protein or polypeptide into a chloroplast or mitochondria of said stamen cells; said fifth DNA being in the same transcriptional unit as said male-sterility DNA and said third promoter and between said male-sterility DNA and said third promoter; and/or

(g) a sixth DNA encoding a transit peptide capable of transporting said fourth protein or polypeptide into a chloroplast or mitochondria of at least said specific tissue or specific cells; said sixth DNA being in the same transcriptional unit as said second marker DNA and said fourth promoter and between said second marker DNA and said fourth promoter.

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48. A pair of parent plants for producing seeds comprising: a) a female-sterile parent plant containing the DNA of any one of claims 1 to 23 incorporated in the nuclear DNA of all of its cells, and b) a female-fertile parent plant.

49. The pair of claim 48 wherein said female-sterile parent plant and said female-fertile parent plant belong to different lines.

50. The pair of claim 48 wherein said female-sterile parent plant and said female-fertile parent plant are derived from the same inbred line.

51. The pair of any one of claims 48 to 50 wherein said female-fertile parent plant is male-sterile.

Patentansprüche

1. DNA, die eine erste chimäre DNA enthält, welche umfasst:

(a) eine DNA für weibliche Sterilität, die eine erste RNA, ein erstes Protein oder Polypeptid kodiert, die bei Erzeugung in Zellen eines weiblichen Fortpflanzungsorgans der Pflanze fähig sind, signifikant den Stoffwechsel, die Funktion und/oder die Entwicklung des weiblichen Fortpflanzungsorgans zu stören; und

(b) einen ersten Promotor, der zur selektiven Ausrichtung der Expression dieser DNA in Zellen der weiblichen Fortpflanzungsorgane der Pflanze fähig ist, wobei die DNA für weibliche Sterilität in der gleichen Transkriptionseinheit wie und unter der Kontrolle des ersten Promotors ist.

2. DNA gemäss Anspruch 1, worin der erste Promotor zur Ausrichtung der Expression der DNA für weibliche Sterilität speziell in Griffelzellen, Narbenzellen, Fruchtknotenzellen, Samenanlagenzellen und/oder Septumzellen fähig ist.

3. DNA gemäss Anspruch 1, worin der erste Promotor zur Ausrichtung der Expression der DNA für weibliche Sterilität speziell in Griffel- und/oder Narbenzellen fähig ist.

4. DNA gemäss einem der Ansprüche 1 bis 3, worin der erste Promotor ausgewählt ist aus der Gruppe, bestehend aus: PSTMG07, das der Promotor eines endogenen Tabakgens ist, das eine mRNA kodiert, aus der die cDNA der Fig. 1A hergestellt werden kann; PSTMG08, das der Promotor eines endogenen Tabakgens ist, das eine mRNA kodiert, aus der die cDNA der Fig. 1B hergestellt werden kann; PSTMG4B12, das der Promotor eines endogenen Tabakgens ist, das eine mRNA kodiert, aus der die cDNA der Fig. 2A hergestellt werden kann; PSTMG3C9, das der Promotor eines endogenen Tabakgens ist, das eine mRNA kodiert, aus der die cDNA der Fig. 2B hergestellt werden kann.

5. DNA gemäss einem der Ansprüche 1 bis 4, worin der erste Promotor ein Promotor eines Gens ist, das dem cDNA-Klon pMON9608 entspricht.

6. DNA gemäss einem der Ansprüche 1 bis 5, worin die DNA für weibliche Sterilität eine Ribonuklease kodiert, insbesondere RNase T1.

7. DNA gemäss einem der Ansprüche 1 bis 5, worin die DNA für weibliche Sterilität eine Barnase kodiert.

8. DNA gemäss einem der Ansprüche 1 bis 5, worin die DNA für weibliche Sterilität eine DNase kodiert, speziell eine Endonuklease, insbesondere EcoRI; eine Protease, speziell ein Papain, insbesondere Papain-Zymogen oder Papain-aktives Protein; eine Glucanase; eine Lipase, insbesondere Phospholipase A₂; eine Lipidperoxidase; einen Zellwandinhibitor; ein Bakteriotoxin; oder ein Ribozym; insbesondere ein Ribozym gegen mRNA, die durch eines der Gene vom STMG-Typ oder durch ein dem cDNA-Klon pMON9608 entsprechendes Gen kodiert wird; oder worin die DNA für weibliche Sterilität alternativ eine Ribonuklease kodiert, insbesondere T₂ oder Rh; ein Glycoprotein kodiert, das durch die S1-, S2-, S3-, S6- oder S7-Allele kodiert wird, insbesondere von Nicotiana glauca; oder eine Antisense-DNA ist, insbesondere eine Antisense-DNA, die eine RNA kodiert, die komplementär zur mRNA eines Gens vom STMG-Typ ist.

9. DNA gemäss einem der Ansprüche 1 bis 5, worin die DNA für weibliche Sterilität ein Enzym kodiert, das die Synthese eines Phytohormons katalysiert, insbesondere das Enzym, das durch Gen 4 von Agrobacterium T-DNA kodiert wird, oder ein Enzym, das durch Gen 1 und/oder Gen 2 von Agrobacterium T-DNA kodiert wird.

10. DNA gemäss einem der Ansprüche 1 bis 5, worin die DNA für weibliche Sterilität eine virusabhängige RNA-Polymerase, insbesondere ein TNV-Replikase, unter der Kontrolle des ersten Promotors kodiert; und ein negatives

strang eines Gens, das das erste Protein oder Polypeptid kodiert; wobei das Gen in der gleichen Transkriptionseinheit wie und unter der Kontrolle eines Promotors ist, wie eines anderen ersten Promotors, der zur Ausrichtung der Expression des Negativstrangs in der Zelle fähig ist; wobei das Gen ebenfalls an seinem 3'-Ende an einen viralen subgenomischen Promotor fusioniert ist, wie den TNV-Subgenompromotor, der spezifisch durch die virusabhängige RNA-Polymerase erkannt wird.

11. DNA gemäss einem der Ansprüche 1 bis 10, worin die DNA ausserdem umfasst:

(e) eine erste DNA, die ein Transitpeptid kodiert, das zum Transport des ersten Proteins oder Polypeptids in einen Chloroplasten oder in Mitochondrien der Zellen des weiblichen Fortpflanzungsorgans fähig ist; wobei die erste DNA in der gleichen Transkriptionseinheit wie die DNA für weibliche Sterilität und der erste Promotor ist und zwischen der DNA für weibliche Sterilität und dem ersten Promotor ist.

12. DNA gemäss einem der Ansprüche 1 bis 10, worin die DNA ausserdem umfasst:

(e) eine dritte DNA, die ein Sekretionssignalpeptid kodiert, das zur Sekretion des ersten Proteins oder Polypeptids ausserhalb der Zelle eines weiblichen Fortpflanzungsorgans einer Pflanze fähig ist; wobei die dritte DNA in der gleichen Transkriptionseinheit wie die DNA für weibliche Sterilität und der erste Promotor ist und sich zwischen der DNA für weibliche Sterilität und dem ersten Promotor befindet.

13. DNA gemäss einem der Ansprüche 1 bis 12, die ebenfalls eine zweite chimäre DNA umfasst, die an die erste chimäre DNA angrenzt, wobei die zweite chimäre DNA umfasst:

(c) eine Marker-DNA, die eine zweite RNA, ein zweites Protein oder Polypeptid kodiert, welche bei Vorliegen wenigstens in einem spezifischen Gewebe oder in wenigstens spezifischen Zellen einer Pflanze die Pflanze leicht abtrennbar von anderen Pflanzen machen, die nicht die zweite RNA, das zweite Protein oder Polypeptid im spezifischen Gewebe oder in den spezifischen Zellen enthalten; und

(d) einen zweiten Promotor, der zur Ausrichtung der Expression der Marker-DNA wenigstens in dem spezifischen Gewebe oder in den spezifischen Zellen fähig ist; wobei die Marker-DNA in der gleichen Transkriptionseinheit wie und unter der Kontrolle des zweiten Promotors ist.

14. DNA gemäss Anspruch 13, worin die Marker-DNA ein Protein kodiert, das die Wirkung eines Herbizids hemmt oder neutralisiert.

15. DNA gemäss Anspruch 14, worin die Marker-DNA ein Herbizidresistenz-Gen ist, insbesondere ein Gen, das Resistenz gegen einen Glutaminsynthetase-Inhibitor wie Phosphinothricin verleiht.

16. DNA gemäss Anspruch 15, worin die Marker-DNA ein *sfrv*- oder *sfrv*-Gen ist.

17. DNA gemäss Anspruch 14, worin die Marker-DNA ein Gen ist, das ein modifiziertes Zielenzym für ein Herbizid mit geringerer Affinität für das Herbizid kodiert, insbesondere eine modifizierte 5-Enolpyruvylshikimat-3-phosphatsynthetase als Ziel für Glyphosat oder eine modifizierte Glutaminsynthetase als Ziel für einen Glutaminsynthetase-Inhibitor wie Phosphinothricin.

18. DNA gemäss Anspruch 13, worin die Marker-DNA ein Gen ist, das ein Protein oder Polypeptid kodiert, das wenigstens dem spezifischen Gewebe oder den spezifischen Zellen eine Färbung verleiht, insbesondere das Gen A1 oder das GUS-Gen; ein Gen, das ein Protein oder Polypeptid kodiert, das der Pflanze Stresstoleranz verleiht, insbesondere das Gen, das Mn-Superoxiddismutase kodiert; oder ein Gen, das ein Protein oder Polypeptid kodiert, das eine Krankheits- oder Schädlingsresistenz verleiht, insbesondere ein Gen, das ein *Bacillus thuringiensis*-Endotoxin kodiert, welches Insektenresistenz verleiht, oder ein Gen, das ein bakterizides Peptid kodiert, das Bakterienresistenz verleiht.

19. DNA gemäss einem der Ansprüche 13 bis 18, worin der zweite Promotor ein konstitutiver Promotor, insbesondere ein PNOS-Promotor oder ein POCS-Promotor ist; ein wundeninduzierbarer Promotor, insbesondere ein TR1'- oder TR2'-Promotor; ein Promotor, der die Genexpression selektiv in Pflanzengewebe mit Photosyntheseaktivität ausrichtet, oder ein Promotor, der die Genexpression selektiv in Blattzellen, Blütenblattzellen oder Samenzellen ausrichtet, insbesondere Samenschalenzellen.

20. DNA gemäss Anspruch 19, worin der zweite Promotor ein 35S-Promotor oder ein SSU-Promotor ist.

21. DNA gemäss einem der Ansprüche 13 bis 20, worin die DNA ausserdem umfasst:

5 (h) eine zweite DNA, die ein Transitpeptid kodiert, das zum Transport des zweiten Proteins oder Polypeptids in einen Chloroplasten oder in Mitochondrien wenigstens des spezifischen Gewebes oder d r spezifischen Zellen fähig ist; wobei die zweite DNA in der gleichen Transkriptionseinheit wie die Marker-DNA und der zweite Promotor ist und zwischen der Marker-DNA und dem zweiten Promotor ist.

10 22. DNA gemäss einem der Ansprüche 13 bis 20, worin die DNA ausserdem umfasst:

(f) eine vierte DNA, die ein Sekretionssignalpeptid kodiert, das zur Sekretion des zweiten Proteins oder Polypeptids ausserhalb wenigstens des spezifischen Gewebes oder spezifischer Zellen fähig ist; wobei die vierte DNA in der gleichen Transkriptionseinheit wie die Marker-DNA und der zweite Promotor ist und sich zwischen der Marker-DNA und dem zweiten Promotor befindet.

23. DNA, die die T-DNA von pMG100 aus Fig. 3, pMG101 aus Fig. 4, pMG102 aus Fig. 5, pMG103 aus Fig. 6, pMG104 aus Fig. 7 oder pMG105 aus Fig. 8 umfasst.

20 24. DNA gemäss einem der Ansprüche 1 bis 23, die eine Zellkern-DNA einer Zelle einer Pflanze oder eines Samens ist.

25. Vektor, der die DNA gemäss einem der Ansprüche 1 bis 23 enthält.

25 26. Zelle einer Pflanze, die die DNA gemäss einem der Ansprüche 1 bis 24 enthält.

27. Zelle einer Pflanze, die stabil in ihre Zellkern-DNA integriert die DNA gemäss einem der Ansprüche 1 bis 23 enthält.

30 28. Zelle gemäss Anspruch 26 oder 27, die zu einer Pflanze regeneriert werden kann, die weiblich-steril ist.

29. Pflanzenzellkultur, die Pflanzenzellen gemäss einem der Ansprüche 26 bis 28 enthält.

30. Pflanze, die die DNA gemäss einem der Ansprüche 1 bis 24 enthält.

35 31. Pflanze, die die DNA gemäss einem der Ansprüche 1 bis 24 in allen ihren Zellen enthält.

32. Pflanze gemäss Anspruch 30 oder 31, die weiblichsteril ist.

40 33. Pflanze gemäss einem der Ansprüche 30 bis 32, die eine Hybridpflanze ist.

34. Weibliche-sterile Pflanze, die eine in das Kerngenom ihrer Zellen aufgenommene Fremd-DNA enthält, worin die Fremd-DNA umfasst:

45 (a) eine DNA für weibliche Sterilität, die eine erste RNA, ein erstes Protein oder Polypeptid kodiert, die bei Erzeugung in Zellen eines weiblichen Fortpflanzungsorgans der Pflanze zur signifikanten Störung des Stoffwechsels, der Funktion und/oder der Entwicklung des weiblichen Fortpflanzungsorgans fähig sind; und

50 (b) einen ersten Promotor, der zur selektiven Ausrichtung der Expression dieser DNA in Zellen der weiblichen Fortpflanzungsorgane der Pflanze fähig ist, wobei die DNA für weibliche Sterilität in der gleichen Transkriptionseinheit wie und unter der Kontrolle des ersten Promotors ist,

mit der Massgabe, dass das Kerngenom der Pflanze homozygot ist, falls der erste Promotor ein Promotor ist, der zur selektiven Ausrichtung der Expression der DNA für weibliche Sterilität in weiblichen Gameten oder in aus weiblichen Gameten stammenden Zellen fähig ist.

35. Pflanze gemäss einem der Ansprüche 30 bis 34, die ausgewählt ist aus der Gruppe bestehend aus Luzerne und Tabak.

36. Pflanzensamen, die die DNA gemäß einem der Ansprüche 1 bis 24 enthält.
37. Samenlose Frucht einer Pflanze, die die DNA gemäß einem der Ansprüche 1 bis 24 enthält.
- 5 38. Verfahren zur Erzeugung einer weiblich-sterilen Pflanze und von Fortpflanzungsmaterial für die Pflanze, welches umfasst:
- (a) Einführen der Fremd-DNA gemäß einem der Ansprüche 1 bis 23 in das Kerngenom einer Pflanzenzelle zum Erhalt einer transformierten Pflanzenzelle;
- 10 (b) Regenerieren der weiblich-sterilen Pflanze aus der transformierten Pflanzenzelle und gegebenenfalls
- (c) Erhalten des Fortpflanzungsmaterials aus der weiblich-sterilen Pflanze, das die Fremd-DNA enthält.
- 15 39. Verfahren gemäß Anspruch 20, worin das Fortpflanzungsmaterial ein Samen ist, insbesondere ein Hybridsamen, und welches ausserdem umfasst:
- (a) Bereitstellen einer Pflanze, die die DNA gemäß einem der Ansprüche 1 bis 23 aufgenommen in die Zellkern-DNA aller ihrer Zellen enthält, und die weiblich-steril ist;
- 20 (b) Fremdbestäuben (i) der weiblich-sterilen Pflanze und (ii) einer weiblich-fruchtbaren Pflanze und
- (c) Gewinnen des Samens aus der weiblichfruchtbaren Pflanze.
- 25 40. Verfahren gemäß Anspruch 20, worin das Fortpflanzungsmaterial ein Samen ist, insbesondere ein Hybridsamen, welches die Schritte umfasst:
- (a) Bereitstellen einer Pflanze, die die DNA gemäß einem der Ansprüche 13 bis 23 enthält, einschliesslich sowohl des zweiten Promotors als auch der Marker-DNA, aufgenommen in die Zellkern-DNA aller ihrer Zellen, und die weiblich-steril ist;
- 30 (b) Fremdbestäuben (i) der weiblich-sterilen Pflanze und (ii) einer weiblich-fruchtbaren Pflanze ohne die Marker-DNA und/oder den zweiten Promotor, und
- 35 (c) Gewinnen des Samens aus der weiblichfruchtbaren Pflanze,
- worin das Verfahren gegebenenfalls die Entfernung ungewünschter weiblich-fruchtbaren Pflanzen auf der Grundlage des Fehlens der Expression der Marker-DNA in den weiblich-fruchtbaren Pflanzen umfasst.
- 40 41. Verfahren zur Erzeugung eines Samens, das die Schritte umfasst aus:
- (a) Bereitstellen einer Pflanze, die die DNA gemäß einem der Ansprüche 13 bis 23 aufgenommen in die Zellkern-DNA aller ihrer Zellen enthält, die weiblich-steril ist, und worin die Marker-DNA ein Herbizidresistenzgen ist, insbesondere ein Gen, das Resistenz gegen einen Glutaminsynthetase-Inhibitor verleiht, wie das sfr- oder sfrv-Gen, oder ein Gen, das ein modifiziertes Zielenzym für ein Herbizid mit geringerer Affinität für ein Herbizid kodiert, insbesondere eine modifizierte 5-Enolpyruvylshikimate-3-phosphatsynthetase als Ziel für Glyphosat oder eine modifizierte Glutaminsynthetase als Ziel für einen Glutaminsynthetase-Inhibitor wie Phosphinothricin,
- 45 (b) Fremdbestäuben (i) der weiblich-sterilen Pflanze und (ii) einer weiblich-fruchtbaren Pflanze ohne die Marker-DNA und/oder den zweiten Promotor, und
- 50 (c) Gewinnen des Samens aus der weiblichfruchtbaren Pflanze,
- 55 worin das Verfahren die Entfernung ungewünschter weiblich-fruchtbaren Pflanzen durch Anwendung des Herbizids umfasst, insbesondere eines Herbizids, das Phosphinothricin umfasst.
42. Verfahren gemäß Anspruch 41, worin der Samen ein Hybridsamen ist.

43. Verfahren gemäss einem der Ansprüche 41 oder 42, worin die weiblich-sterile Pflanze zusätzlich zum Herbizidresistenzgen wenigstens eine andere, stabil in das Kerngenom ihrer Zellen integrierte Marker-DNA im gleichen genetischen Locus wie die DNA für weibliche Sterilität enthält und die weiblichfruchtbare Pflanze die andere Marker-DNA nicht enthält.

44. Verfahren gemäss einem der Ansprüche 39 bis 43, worin die weiblich-fruchtbare Pflanze männlich-steril ist.

45. Verfahren gemäss Anspruch 44, worin die weiblichfruchtbare, männlich-sterile Pflanze eine in das Kerngenom ihrer Zellen aufgenommene Fremd-DNA umfasst, worin die Fremd-DNA umfasst:

(a) eine DNA für männliche Sterilität, die eine dritte RNA, ein drittes Protein oder Polypeptid kodiert, welche bei Erzeugung in Staubblattzellen der Pflanze zur signifikanten Störung des Stoffwechsels, der Funktion und/oder der Entwicklung der Staubblattzellen fähig sind;

(b) einen dritten Promotor, der zur selektiven Ausrichtung der Expression dieser DNA in Staubblattzellen der Pflanze fähig ist, wobei die DNA für männliche Sterilität in der gleichen Transkriptionseinheit wie und unter der Kontrolle des dritten Promotors ist;

mit der Massgabe, dass das Kerngenom der transformierten Pflanzenzelle homozygot ist, falls der dritte Promotor ein Promotor ist, der zur selektiven Ausrichtung der Expression der DNA für männliche Sterilität in Pollenzellen fähig ist.

46. Verfahren gemäss Anspruch 45, worin die Fremd-DNA der weiblich-fruchtbaren, männlich-sterilen Pflanze ebenfalls umfasst:

(d) eine zweite Marker-DNA, die eine vierte RNA, ein viertes Protein oder Polypeptid kodiert, welche bei Vorliegen wenigstens in einem spezifischen Gewebe oder in wenigstens spezifischen Zellen der Pflanze leicht abtrennbar von anderen Pflanzen machen, die die vierte RNA, das vierte Protein oder Polypeptid im spezifischen Gewebe oder in den spezifischen Zellen nicht enthalten; und

(e) einen vierten Promotor, der zur Ausrichtung der Expression der zweiten Marker-DNA wenigstens im spezifischen Gewebe oder in den spezifischen Zellen fähig ist; wobei die zweite Marker-DNA in der gleichen Transkriptionseinheit wie und unter der Kontrolle des vierten Promotors ist.

47. Verfahren gemäss einem der Ansprüche 45 oder 46, worin die Fremd-DNA ausserdem umfasst:

(f) eine fünfte DNA, die ein Transitpeptid kodiert, das zum Transport des dritten Proteins oder Polypeptids in einen Chloroplasten oder in Mitochondrien der Staubblattzellen fähig ist; wobei die fünfte DNA in der gleichen Transkriptionseinheit wie die DNA für männliche Sterilität und der dritte Promotor ist und zwischen der DNA für männliche Sterilität und dem dritten Promotor ist; und/oder

(g) eine sechste DNA, die ein Transitpeptid kodiert, das zum Transport des vierten Proteins oder Polypeptids in einen Chloroplasten oder in Mitochondrien wenigstens des spezifischen Gewebes oder der spezifischen Zellen fähig ist; wobei die sechste DNA in der gleichen Transkriptionseinheit wie die zweite Marker-DNA und der vierte Promotor ist und zwischen der zweiten Marker-DNA und dem vierten Promotor ist.

48. Paar von Mutterpflanzen zur Erzeugung von Samen, umfassend:

(a) eine weiblich-sterile Mutterpflanze, die die DNA gemäss einem der Ansprüche 1 bis 23 aufgenommen in die Zellkern-DNA aller ihrer Zellen enthält, und

(b) eine weiblich-fruchtbare Mutterpflanze.

49. Paar gemäss Anspruch 48, worin die weiblich-sterile Mutterpflanze und die weiblich-fruchtbare Mutterpflanze zu unterschiedlichen Linien gehören.

50. Paar gemäss Anspruch 48, worin die weiblich-sterile Mutterpflanze und die weiblich-fruchtbare Mutterpflanze aus der gleichen Inzuchtlinie stammen.

51. Paar gemäss einem der Ansprüche 48 bis 50, worin die weiblich-fruchtbar Mutterpflanze männlich-steril ist.

R revendications

- 5 1. ADN qui contient un premier ADN chimérique qui comprend :
 - (a) un ADN de stérilité femelle codant pour un premier ARN, une protéine ou un polypeptide, capable, quand il est produit dans des cellules d'un organe de reproduction femelle de ladite plante de perturber significativement le métabolisme, le fonctionnement et/ou le développement de l'organe de reproduction femelle ; et
 - 10 (b) un premier promoteur capable de diriger l'expression de cet ADN sélectivement dans des cellules des organes de reproduction femelle de ladite plante, ledit ADN de stérilité femelle étant dans la même unité transcriptionnelle que, et sous la régulation dudit premier promoteur.
- 15 2. ADN selon la revendication 1, dans lequel ledit premier promoteur est capable de diriger l'expression dudit ADN de stérilité femelle en particulier dans des cellules de style, des cellules de stigmate, des cellules d'ovaire, des cellules d'ovule et/ou des cellules de septum.
3. ADN selon la revendication 1, dans lequel ledit premier promoteur est capable de diriger l'expression dudit ADN de stérilité femelle en particulier dans des cellules de style et/ou de stigmate.
- 20 4. ADN selon l'une quelconque des revendications 1 à 3, dans lequel ledit premier promoteur est choisi dans le groupe constitué de : PSTMG07 qui est le promoteur d'un gène endogène de tabac codant pour un ARNm d'où l'ADNc de la figure 1A peut être préparé ; PSTMG08 qui est le promoteur d'un gène endogène de tabac codant pour un ARNm d'où l'ADNc de la figure 1B peut être préparé ; PSTMG4B12 qui est le promoteur d'un gène endogène de tabac codant pour un ARNm d'où l'ADNc de la figure 2A peut être préparé ; PSTMG3C9 qui est le promoteur d'un gène endogène de tabac codant pour un ARNm d'où l'ADNc de la figure 2B peut être préparé.
- 25 5. ADN selon l'une quelconque des revendications 1 à 4, dans lequel ledit premier promoteur est un promoteur d'un gène correspondant à un clone d'ADNc pMON9608.
- 30 6. ADN selon l'une quelconque des revendications 1 à 5, dans lequel ledit ADN de stérilité femelle code pour une ribonucléase, en particulier l'ARNase T1.
7. ADN selon l'une quelconque des revendications 1 à 5, dans lequel ledit ADN de stérilité femelle code pour une BARNase.
- 35 8. ADN selon l'une quelconque des revendications 1 à 5, dans lequel ledit ADN de stérilité femelle code pour une ADNase, en particulier une endonucléase, particulièrement EcoRI ; une protéase, en particulier une papaine, particulièrement le zymogène de papaine ou une protéine active de papaine ; une glucanase ; une lipase, en particulier la phospholipase A₂ ; une peroxydase de lipide ; un inhibiteur de paroi cellulaire ; une toxine bactérienne, ou une ribozyme ; en particulier une ribozyme contre ARNm codé par l'un quelconque des gènes de type STMG, ou par un gène correspondant au clone ADNc pMON9608 ; ou en variante, dans lequel ledit ADN de stérilité femelle : code pour une ribonucléase, en particulier T₂ ou Rh ; code pour une glycoprotéine codée par les allèles S1, S2, S3, S6 ou S7, en particulier de Nicotiana glauca ; ou est un ADN anti-sens, en particulier un ADN anti-sens codant pour un ARN qui est complémentaire à l'ARNm d'un gène de type STMG.
- 45 9. ADN selon l'une quelconque des revendications 1 à 5, dans lequel ledit ADN de stérilité femelle code pour une enzyme qui catalyse la synthèse d'une phytohormone, en particulier l'enzyme codée par le gène 4 de ADN-T d'Agrobacterium, ou une enzyme codée par le gène 1 et/ou le gène 2 d'ADN-T d'Agrobacterium.
- 50 10. ADN selon l'une quelconque des revendications 1 à 5, dans lequel ledit ADN de stérilité femelle code pour une polymérase d'ARN dépendante d'un virus, en particulier une TNV répliquase, sous la régulation dudit premier promoteur ; et un brin négatif d'un gène codant pour ledit premier polypeptide ou protéine ; ledit gène étant dans la même unité transcriptionnelle que, et sous la régulation de, un promoteur, tel qu'un autre premier promoteur, capable de diriger l'expression dudit brin négatif dans la cellule ; ledit gène étant également fusionné à son extrémité 3' à un promoteur subgénomique viral, tel que le promoteur TNV subgénomique , qui est spécifiquement reconnu par ladite polymérase d'ARN dépendante de virus.
- 55

11. ADN selon l'une quelconque des revendications 1 à 10, dans lequel ledit ADN comprend de plus :

(e) un premier ADN codant pour un peptide de transit capable de transporter ledit premier polypeptide ou protéine dans un chloroplaste ou une mitochondrie desdites cellules d'organe de reproduction femelle ; ledit premier ADN étant dans la même unité transcriptionnelle que ledit ADN de stérilité femelle et ledit premier promoteur, et situé ntr ledit ADN de stérilité femelle et ledit premier promoteur.

12. ADN selon l'une quelconque des revendications 1 à 10, dans lequel ledit ADN comprend de plus :

(e) un troisième ADN codant pour un peptide signal sécrétoire capable de sécréter ledit premier poly-peptide ou protéine à l'extérieur de la cellule d'un organe de reproduction femelle d'une plante ; ledit troisième ADN étant dans la même unité transcriptionnelle que ledit ADN de stérilité femelle et ledit premier promoteur, et situé entre ledit ADN de stérilité femelle et ledit premier promoteur.

13. ADN selon l'une quelconque des revendications 1 à 12, qui comprend également un deuxième ADN chimérique, adjacent audit premier ADN chimérique, ledit deuxième ADN chimérique comprenant :

(c) un ADN marqueur codant pour un deuxième ARN, polypeptide ou protéine qui, quand il est présent au moins dans un tissu spécifique ou dans au moins des cellules spécifiques d'une plante, rend ladite plante facilement séparable des autres plantes qui ne contiennent pas ledit second ARN, polypeptide ou protéine dans ledit tissu spécifique ou lesdites cellules spécifiques ; et

(d) un deuxième promoteur capable de diriger l'expression dudit ADN marqueur au moins dans ledit tissu spécifique ou lesdites cellules spécifiques ; ledit ADN marqueur étant dans la même unité transcriptionnelle que ledit, et sous la régulation dudit, deuxième promoteur.

14. ADN selon la revendication 13, dans lequel ledit ADN marqueur code pour une protéine inhibant ou neutralisant l'action d'un herbicide.

15. ADN selon la revendication 14, dans lequel ledit ADN marqueur est un gène de résistance à un herbicide, en particulier un gène conférant une résistance à un inhibiteur de glutamine synthétase, tel que la phosphinothricine.

16. ADN selon la revendication 15, dans lequel ledit ADN marqueur est un gène sfr ou sfrv.

17. ADN selon la revendication 14, dans lequel ledit ADN marqueur est un gène codant pour une enzyme cible modifiée pour un herbicide ayant une affinité inférieure pour l'herbicide, en particulier une 5-énolpyruvylshikimate-3 phosphate synthase modifiée en tant que cible pour un glyphosate ou une glutamine synthétase modifiée en tant que cible pour un inhibiteur de glutamine synthétase tel que la phosphinothricine.

18. ADN selon la revendication 13, dans lequel ledit ADN marqueur est un gène codant pour une protéine ou un polypeptide conférant une couleur à au moins ledit tissu spécifique ou lesdites cellules spécifiques, en particulier le gène A1 ou le gène GUS ; un gène codant pour une protéine ou un polypeptide conférant une tolérance au stress à ladite plante, en particulier le gène codant pour la Mn-superoxyde dismutase ; ou un gène codant pour une protéine ou un polypeptide conférant une résistance aux maladies ou aux nuisibles, en particulier un gène codant pour une endotoxine de *Bacillus thuringiensis* qui confère une résistance aux insectes ou un gène codant pour un peptide bactéricide qui confère une résistance bactérienne.

19. ADN selon l'une quelconque des revendications 13 à 18, dans lequel ledit deuxième promoteur est : un promoteur constitutif, en particulier un promoteur PNOS ou un promoteur POCS ; un promoteur inductible par blessure, en particulier un promoteur TR1' ou TR2' ; un promoteur qui dirige l'expression d'un gène sélectivement dans un tissu de plante ayant une activité de photosynthèse, ou un promoteur qui dirige l'expression d'un gène sélectivement dans des cellules de feuille, des cellules de pétale ou des cellules de graine, en particulier des cellules d'enveloppe de graine.

20. ADN selon la revendication 19, dans lequel ledit deuxième promoteur est un promoteur 35S ou un promoteur SSU.

21. ADN selon l'une quelconque des revendications 13 à 20, dans lequel ledit ADN comprend de plus :

(h) un deuxième ADN codant pour un peptide de transit capable de transporter ledit deuxième polypeptide ou

protéine dans un chloroplaste ou une mitochondrie d'au moins ledit tissu spécifique ou lesdites cellules spécifiques ; ledit deuxième ADN étant dans la même unité transcriptionnelle que ledit ADN marqueur et ledit deuxième promoteur, et situé entre ledit ADN marqueur et ledit deuxième promoteur.

- 5 22. ADN selon l'une quelconque des revendications 13 à 20, dans lequel ledit ADN comprend de plus :

10 (f) un quatrième ADN codant pour un peptide signal de sécrétion capable de sécréter ledit deuxième polypeptide ou protéine à l'extérieur d'au moins le tissu spécifique ou les cellules spécifiques ; ledit quatrième ADN étant dans la même unité transcriptionnelle que ledit ADN marqueur et ledit deuxième promoteur, et situé entre ledit ADN marqueur et ledit deuxième promoteur.
23. ADN qui comprend l'ADN-T de pMG100 de la figure 3, pMG101 de la figure 4, pMG102 de la figure 5, pMG103 de la figure 6, pMG104 de la figure 7, ou pMG105 de la figure 8.
- 15 24. ADN selon l'une quelconque des revendications 1 à 23 qui est un ADN nucléaire d'une cellule d'une plante ou d'une graine.
25. Vecteur qui contiennent l'ADN selon l'une quelconque des revendications 1 à 23.
- 20 26. Cellule d'une plante qui contient l'ADN selon l'une quelconque des revendications 1 à 24.
27. Cellule d'une plante qui contient, intégré de façon stable dans son ADN nucléaire, l'ADN selon l'une quelconque des revendications 1 à 23.
- 25 28. Cellule selon la revendication 26 ou la revendication 27 qui peut être régénérée dans une plante qui est stérile femelle.
29. Culture de cellule végétale contenant des cellules végétales selon l'une quelconque des revendications 26 à 28.
- 30 30. Plante qui contient l'ADN selon l'une quelconque des revendications 1 à 24.
31. Plante qui contient l'ADN selon l'une quelconque des revendications 1 à 24 dans la totalité de ses cellules.
32. Plante selon la revendication 30 ou 31 qui est stérile femelle.
- 35 33. Plante selon l'une quelconque des revendications 30 à 32 qui est une plante hybride.
34. Plante stérile femelle contenant un ADN étranger incorporé dans le génome nucléaire de ses cellules, dans laquelle ledit ADN étranger comprend :

40 (a) un ADN de stérilité femelle codant pour un premier ARN, protéine ou polypeptide, capable quand il est produit dans des cellules d'un organe de reproduction femelle de ladite plante, de perturber de façon significative le métabolisme, le fonctionnement et/ou le développement de l'organe de reproduction femelle ; et
45 (b) un premier promoteur capable de diriger l'expression de cet ADN sélectivement dans des cellules des organes de reproduction femelle de ladite plante, ledit ADN de stérilité femelle étant dans la même unité transcriptionnelle que ledit, et sous la régulation dudit, premier promoteur,

à condition que, si ledit premier promoteur est un promoteur capable de diriger l'expression dudit ADN de stérilité femelle sélectivement dans des gamètes femelles, ou dans des cellules provenant de gamètes femelles, le
50 génome nucléaire de ladite plante est homozygote.
35. Plante selon l'une quelconque des revendications 30 à 34 qui est choisie dans le groupe formé de la luzerne et du tabac.
- 55 36. Graine végétale qui contient l'ADN selon l'une quelconque des revendications 1 à 24.
37. Fruit sans graine d'une plante qui contient l'ADN selon l'une quelconque des revendications 1 à 24.

38. Procédé pour produire une plante stérile femelle et un matériel de reproduction pour ladite plante, qui comprend :

- a) l'introduction d'un ADN étranger selon l'une quelconque des revendications 1 à 23 dans le génome nucléaire d'une cellule végétale afin d'obtenir une cellule végétale transformée ;
- b) la régénération de ladite plante stérile femelle à partir de ladite cellule végétale transformée et ; éventuellement,
- c) l'obtention dudit matériel de reproduction à partir de ladite plante stérile femelle qui comprend ledit ADN étranger.

39. Procédé selon la revendication 20, dans lequel ledit matériel de reproduction est une graine, en particulier une graine hybride, et qui comprend de plus :

- a) la fourniture d'une plante qui contient l'ADN selon l'une quelconque des revendications 1 à 23 incorporé dans l'ADN nucléaire de la totalité de ses cellules et qui est stérile femelle ;
- b) la pollinisation croisée i) de ladite plante stérile femelle, et ii) d'une plante fertile femelle, et
- c) la récupération de ladite graine de ladite plante fertile femelle.

40. Procédé selon la revendication 20, dans lequel ledit matériel de reproduction est une graine, en particulier une graine hybride, qui comprend les étapes consistant :

- a) à fournir une plante qui contient l'ADN selon l'une quelconque des revendications 13 à 23, comprenant à la fois ledit deuxième promoteur et ledit ADN marqueur, incorporé dans l'ADN nucléaire de la totalité de ses cellules, et qui est stérile femelle ;
- b) à réaliser une pollinisation croisée i) de ladite plante stérile femelle, et ii) une plante fertile femelle sans ledit ADN marqueur et/ou ledit deuxième promoteur, et
- c) à récupérer la graine à partir de ladite plante fertile femelle,

dans lequel ledit procédé comprend éventuellement l'élimination des plantes fertiles femelles non souhaitables sur la base de l'absence d'expression de l'ADN marqueur dans lesdites plantes fertiles femelles.

41. Procédé pour produire une graine, qui comprend les étapes consistant :

- (a) à fournir une plante qui contient l'ADN selon l'une quelconque des revendications 13 à 23 incorporé dans l'ADN nucléaire de la totalité de ses cellules, qui est stérile femelle, et dans lequel ledit ADN marqueur est un gène de résistance à un herbicide, en particulier un gène conférant une résistance à un inhibiteur de glutamine synthétase tel que le gène sfr ou sfrv, ou un gène codant pour une enzyme cible modifiée pour un herbicide ayant une affinité inférieure pour un herbicide, en particulier une 5-énoypyruvylshikimate-3 phosphate synthase modifiée en tant que cible pour un glyphosate ou une glutamine synthétase modifiée comme cible pour un inhibiteur de glutamine synthase tel que la phosphinothricine,
- (b) à réaliser une pollinisation croisée i) de ladite plante stérile femelle, et ii) d'une plante fertile femelle sans ledit ADN marqueur et/ou ledit deuxième promoteur, et
- (c) à récupérer la graine provenant de ladite plante fertile femelle,

dans lequel ledit procédé comprend l'élimination des plantes fertiles femelles non souhaitables par l'application dudit herbicide, en particulier un herbicide contenant de la phosphinothricine.

42. Procédé selon la revendication 41, dans lequel ladite graine est une graine hybride.

43. Procédé selon l'une quelconque des revendications 41 ou 42, dans lequel ladite plante stérile femelle contient en plus dudit gène de résistance à un herbicide au moins un autre ADN marqueur intégré de façon stable dans le génome nucléaire de ses cellules dans le même locus génétique que ledit ADN de stérilité femelle et ladite plante fertile femelle ne contient pas ledit autre ADN marqueur.

44. Procédé selon l'une quelconque des revendications 39 et 43, dans lequel la plante fertile femelle est stérile mâle.

45. Procédé selon la revendication 44, dans lequel ladite plante fertile femelle et stérile mâle comprend un ADN étranger incorporé dans le génome nucléaire de ses cellules, dans lequel ledit ADN étranger comprend :

(a) un ADN de stérilité mâle codant pour un troisième ARN, protéine ou polypeptide, capable quand il est produit dans des cellules d'étamine de ladite plante, de perturber significativement le métabolisme, le fonctionnement et/ou le développement des cellules d'étamine ;

(b) un troisième promoteur capable de diriger l'expression de cet ADN sélectivement dans des cellules d'étamine de ladite plante, ledit ADN de stérilité mâle étant dans la même unité transcriptionnelle que ledit, et sous la régulation dudit, troisième promoteur ;

à condition que, si ledit troisième promoteur est un promoteur capable de diriger l'expression dudit ADN de stérilité mâle sélectivement dans des cellules de pollen, le génome nucléaire de ladite cellule végétale transformée est homozygote.

46. Procédé selon la revendication 45, dans lequel ledit ADN étranger de ladite plante fertile femelle stérile mâle comprend également :

(d) un deuxième ADN marqueur codant pour un quatrième ARN, protéine ou polypeptide qui, quand il est présent au moins dans un tissu spécifique ou dans au moins des cellules spécifiques de ladite plante, rend ladite plante facilement séparable des autres plantes qui ne contiennent pas ledit quatrième ARN, protéine ou polypeptide dans ledit tissu spécifique ou lesdites cellules spécifiques ; et

(e) un quatrième promoteur capable de diriger l'expression dudit deuxième ADN marqueur au moins dans ledit tissu spécifique ou lesdites cellules spécifiques ; ledit deuxième ADN marqueur étant dans la même unité transcriptionnelle que ledit, et sous la régulation dudit, quatrième promoteur.

47. Procédé selon l'une quelconque des revendications 45 ou 46, dans lequel ledit ADN étranger comprend de plus :

(f) un cinquième ADN codant pour un peptide de transit afin de transporter ledit troisième polypeptide ou protéine dans un chloroplaste ou une mitochondrie desdites cellules d'étamine ; ledit cinquième ADN étant dans la même unité transcriptionnelle que ledit ADN de stérilité mâle et ledit troisième promoteur, et situé entre ledit ADN de stérilité mâle et ledit troisième promoteur ; et/ou

(g) un sixième ADN codant pour un peptide de transit capable de transporter ledit quatrième polypeptide ou protéine dans un chloroplaste ou une mitochondrie d'au moins ledit tissu spécifique ou lesdites cellules spécifiques ; ledit sixième ADN étant dans la même unité transcriptionnelle que ledit deuxième ADN marqueur et ledit quatrième promoteur, et situé entre ledit deuxième ADN marqueur et ledit quatrième promoteur.

48. Paire de plantes parents pour produire des graines comprenant : a) une plante parent stérile femelle contenant l'ADN selon l'une quelconque des revendications 1 à 23 incorporé dans l'ADN nucléaire de la totalité de ses cellules, et b) une plante parent fertile femelle.

49. Paire de la revendication 48, dans laquelle ladite plante parent stérile femelle et ladite plante parent fertile femelle appartiennent à différentes lignées.

50. Paire de la revendication 48, dans laquelle ladite plante parent stérile femelle et ladite plante parent fertile femelle proviennent de la même lignée consanguine.

51. Paire selon l'une quelconque des revendications 48 à 50, dans laquelle ladite plante parent fertile femelle est stérile mâle.

Figure 1 A

		9			18			27			36	
GAA	TTC	CCA	TTG	CCT	TTC	GAA	TTG	CCA	CCA	GCG	GAG	ATC
Glu	Phe	Pro	Leu	Pro	Phe	Glu	Leu	Pro	Pro	Ala	Glu	Ile
		45			54			63			72	
CCA	TTG	CCG	GAG	ATC	CCA	TTG	CCT	TTC	GAT	GGG	CCT	ACA
Pro	Leu	Pro	Glu	Ile	Pro	Leu	Pro	Phe	Asp	Gly	Pro	Thr
		81			90			99			108	
TTC	GTG	CTA	CCG	CCA	CCA	TCA	CCA	CCA	CCA	CCT	CCA	TCG
Phe	Val	Leu	Pro	Pro	Pro	Ser	Pro	Pro	Pro	Pro	Pro	Ser
		126			135			144			153	
TCA	CCA	TCT	CCA	TCT	CCA	GCA	AAG	CAA	TCA	CCA	CCA	CCT
Ser	Pro	Ser	Pro	Ser	Pro	Ala	Lys	Gln	Ser	Pro	Pro	Pro
		162			171			180			189	
CCT	CGG	GCA	CCA	TCA	CCA	TCA	CCA	GCT	ACT	CAG	CCA	CCT
Pro	Arg	Ala	Pro	Ser	Pro	Ser	Pro	Ala	Thr	Gln	Pro	Pro
		198			207			216			225	
ATA	AAG	CAA	CCG	CCA	CCA	CCA	AGT	GCT	AAG	AAA	TCT	CCT
Ile	Lys	Gln	Pro	Pro	Pro	Pro	Ser	Ala	Lys	Lys	Ser	Pro
		243			252			261			270	
CCG	CCA	CCA	GTT	GCT	TAT	CCA	CCA	GTT	ATG	GCA	CCA	TCT
Pro	Pro	Pro	Val	Ala	Tyr	Pro	Pro	Val	MET	Ala	Pro	Ser
		279			288			297			306	
CCA	TCA	CCG	GCT	GCT	GAG	CCA	CCT	ATT	ATA	GCA	CCA	TTT
Pro	Ser	Pro	Ala	Ala	Glu	Pro	Pro	Ile	Ile	Ala	Pro	Phe

Figure 1 A (cont. 1)

315	324	333	342	351
CCA TCA CCA	ACA GCG AAT	CTA CCC CTT	ATT CCC CGT	CGA
Pro Ser Pro	Thr Ala Asn	Leu Pro Leu	Ile Pro Arg	Arg
360	369	378	387	
CCA GCA CCA	CCA GTA GTT	AAG CCG CTT	CCA CCT TTG	GGG
Pro Ala Pro	Pro Val Val	Lys Pro Leu	Pro Pro Leu	Gly
396	405	414	423	
AAG CCC CCT	ATC GTC AAT	GGC CTT GTT	TAT TGT AAA	TCC
Lys Pro Pro	Ile Val Asn	Gly Leu Val	Tyr Cys Lys	Ser
432	441	450	459	468
TGC AAC AGC	TAT GGG TTC	CCC ACT CTG	CTC AAC ACC	TCC
Cys Asn Ser	Tyr Gly Phe	Pro Thr Leu	Leu Asn Thr	Ser
477	486	495	504	
CTA CTC CCA	GGA GCT GTT	GTG AAA CTA	GTT TGC TAC	AAC
Leu Leu Pro	Gly Ala Val	Val Lys Leu	Val Cys Tyr	Asn
513	522	531	540	
GGA AAG AAA	ACA ATG GTT	CAA TCG GCG	ACG ACA GAC	AAC
Gly Lys Lys	Thr MET Val	Gln Ser Ala	Thr Thr Asp	Asn
549	558	567	576	585
AAA GGT GAG	TTT CGG ATC	ATT CCC AAA	TCA TTA ACC	AGA
Lys Gly Glu	Phe Arg Ile	Ile Pro Lys	Ser Leu Thr	Arg
594	603	612	621	
GCA GAT GTT	GGC AAG TGC	AAG TTA TAT	TTA GTG AAA	TCA
Ala Asp Val	Gly Lys Cys	Lys Leu Tyr	Leu Val Lys	Ser

Figure 1 A (cont. 2)

630	639	648	657	
CCA AAT CCA AAT TGC AAT GTC CCA ACA AAT TTC AAT GGT				
Pro Asn Pro Asn Cys Asn Val Pro Thr Asn Phe Asn Gly				
666	675	684	693	702
GGA AAA TCT GGT GGT TTA TTG AAG CCT CTC CTA CCA CCT				
Gly Lys Ser Gly Gly Leu Leu Lys Pro Leu Leu Pro Pro				
711	720	729	738	
AAA CAA CCG ATT ACC CCT GCC GCT GTC CCT CTA TCT GAT				
Lys Gln Pro Ile Thr Pro Ala Ala Val Pro Leu Ser Asp				
747	756	765	774	
TTA TAT GGT GTT GGA CCT TTT ATA TTT GAA GCC TCC AGC				
Leu Tyr Gly Val Gly Pro Phe Ile Phe Glu Ala Ser Ser				
783	792	801	810	819
AAA ATG CCA TGC GAT AAG AAT TGA GCT CCT CAT TAC TAG				
Lys MET Pro Cys Asp Lys Asn . Ala Pro His Tyr .				
828	837	846	855	
AGC GAT AAT GTA TAA GAG CAT GAG TTT GTG ACG GAA ATT				
Ser Asp Asn Val . Glu His Glu Phe Val Thr Glu Ile				
864	873	882	891	
ATT TTT TTC TTT TTT GTT CTA TAG TTT ATA CAA GGA GAC				
Ile Phe Phe Phe Phe Val Leu . Phe Ile Gln Gly Asp				
900	909	918	927	936
AGA AAA CTT TGT ATC ACT ATA CAG AAA TCA AAT GAG TCG				
Arg Lys Leu Cys Ile Thr Ile Gln Lys Ser Asn Glu Ser				

Figure 1 A (cont. 3)

	945		954		963			
CAA	AAG	TCA	AAA	TCG	AAT	TTA	TGA	AAA
Gln	Lys	Ser	Lys	Ser	Asn	Leu	.	Lys

Figure 1 B

				11					20					29					38					
GA	ATT	CCG	GCT	TTT	ACA	TCA	GTA	AAG	ATC	CTA	GTG	CTC												
Ile	Pro	Ala	Phe	Thr	Ser	Val	Lys	Ile	Leu	Val	Leu													
				47					56					65					74					
ATA	CAA	GTT	TCA	GTT	TTA	GCA	CTC	AGC	TCA	TTC	TCA	GAG												
Ile	Gln	Val	Ser	Val	Leu	Ala	Leu	Ser	Ser	Phe	Ser	Glu												
				83					92					101					110					
CTT	AGC	TTT	GGT	AAA	GGA	ATT	GAA	AGC	TCG	TCA	TTA	GAC												
Leu	Ser	Phe	Gly	Lys	Gly	Ile	Glu	Ser	Ser	Ser	Leu	Asp												
				119					128					137					146					155
AAA	GGA	CAA	CAC	CAT	CCA	ATC	TTC	TCA	ACA	GTT	CAC	TTA												
Lys	Gly	Gln	His	His	Pro	Ile	Phe	Ser	Thr	Val	His	Leu												
				164					173					182					191					
TTC	TTT	GGA	AAG	TCT	CCC	AAG	AAA	AGC	CCC	TCT	AGC	CCT												
Phe	Phe	Gly	Lys	Ser	Pro	Lys	Lys	Ser	Pro	Ser	Ser	Pro												
				200					209					218					227					
ACA	CCG	GTA	AAC	AAG	CCA	TCA	CCA	TCA	CCA	CCA	CCA	CAG												
Thr	Pro	Val	Asn	Lys	Pro	Ser	Pro	Ser	Pro	Pro	Pro	Gln												
				236					245					254					263					272
GTT	AAG	TCA	TCC	CTT	CCG	CCG	CCT	GCT	AAG	TCA	CCA	CCG												
Val	Lys	Ser	Ser	Leu	Pro	Pro	Pro	Ala	Lys	Ser	Pro	Pro												
				281					290					299					308					
CCG	CCA	CCA	GCT	AAG	TCA	CCA	CCT	CCT	CTG	CTG	CCT	CCA												
Pro	Pro	Pro	Ala	Lys	Ser	Pro	Pro	Pro	Leu	Leu	Pro	Pro												

Figure 1 B (cont. 1)

317	326	335	344	
CCA CCA TCT CAA	CCA CCA AAA CAA	CCA CCT CCA CCT	CCG	
Pro Pro Ser Gln	Pro Pro Lys Gln	Pro Pro Pro Pro		
353	362	371	380	389
CCG CCA CCA GCA	AAG CAA CCA CCA	TCT GCT AAG CCA	CCT	
Pro Pro Pro Ala	Lys Gln Pro Pro	Ser Ala Lys Pro	Pro	
398	407	416	425	
ATT AAA CCT CCA	TCT CCG TCA CCG	GCT GCT CAG CCA	CCA	
Ile Lys Pro Pro	Ser Pro Ser Pro	Ala Ala Gln Pro	Pro	
434	443	452	461	
GCA ACG CAA CGA	GCA ACA CCA CCA	CCG GCA ATG CAA	CGG	
Ala Thr Gln Arg	Ala Thr Pro Pro	Pro Ala MET Gln	Arg	
470				
GCA CC				
Ala				

Figure 2 A

	9		18		27		36					
GCC	CTG	TAG	CGG	CAT	TAA	GCG	CGG	CGG	GTG	TGG	TGG	TTA
Ala	Leu	.	Arg	His	.	Ala	Arg	Arg	Val	Trp	Trp	Leu
	45		54		63		72					
CGC	GCA	GTG	ACC	GCT	ACA	CTT	GCC	AGC	GCC	CTA	GCG	CCC
Arg	Ala	Val	Thr	Ala	Thr	Leu	Ala	Ser	Ala	Leu	Ala	Pro
	81		90		99		108		117			
GCT	CCT	TTC	GCT	CTT	CTC	ATT	CTC	ATC	ATC	CTC	ACT	CTT
Ala	Pro	Phe	Ala	Leu	Leu	Ile	Leu	Ile	Ile	Leu	Thr	Leu
	126		135		144		153					
TCT	AGC	ACA	CCA	ATT	ACC	ACA	ATG	TCT	ATA	CCC	GAG	ACA
Ser	Ser	Thr	Pro	Ile	Thr	Thr	MET	Ser	Ile	Pro	Glu	Thr
	162		171		180		189					
AAC	CGT	AGA	AAT	GCA	ACT	ACA	AAC	TCT	TAC	ACC	GAT	GTT
Asn	Arg	Arg	Asn	Ala	Thr	Thr	Asn	Ser	Tyr	Thr	Asp	Val
	198		207		216		225		234			
GCT	CTT	TCT	GCG	CGA	AAA	GGT	GCA	TTT	CCT	CCT	CCC	AGA
Ala	Leu	Ser	Ala	Arg	Lys	Gly	Ala	Phe	Pro	Pro	Pro	Arg
	243		252		261		270					
AAG	CTA	GGA	GAA	TAC	TCG	ACA	AAT	TCT	ACC	GAC	TAC	AAC
Lys	Leu	Gly	Glu	Tyr	Ser	Thr	Asn	Ser	Thr	Asp	Tyr	Asn
	279		288		297		306					
TTG	ATC	TGC	AAA	ACT	TGC	AAG	AGA	TTA	TCG	GAA	CGC	AAT
Leu	Ile	Cys	Lys	Thr	Cys	Lys	Arg	Leu	Ser	Glu	Arg	Asn

Figure 2 A (cont. 1)

315		324		333		342		351				
ACA	TGT	TGT	TTC	AAC	TAC	AGT	TGT	GTT	GAT	GTG	TCC	ACC
Thr	Cys	Cys	Phe	Asn	Tyr	Ser	Cys	Val	Asp	Val	Ser	Thr

	360		369		378		387					
AAC	AGG	TTC	AAC	TGT	GGC	TCC	TGT	GGC	CTT	GTG	TGT	AAC
Asn	Arg	Phe	Asn	Cys	Gly	Ser	Cys	Gly	Leu	Val	Cys	Asn

	396		405		414		423					
CTT	GGA	ACG	AGA	TGC	TGT	GGT	GGG	ATC	TGT	GTG	GAC	ATC
Leu	Gly	Thr	Arg	Cys	Cys	Gly	Gly	Ile	Cys	Val	Asp	Ile

432		441		450		459		468				
CAG	AAA	GAC	AAT	GGC	AAT	TGT	GGC	AAG	TGT	TCT	AAT	GTT
Gln	Lys	Asp	Asn	Gly	Asn	Cys	Gly	Lys	Cys	Ser	Asn	Val

	477		486		495		504					
TGC	TCT	CCT	GGT	CAG	AAG	TGT	TCA	TTT	GGG	TTT	TGT	GAC
Cys	Ser	Pro	Gly	Gln	Lys	Cys	Ser	Phe	Gly	Phe	Cys	Asp

	513		522		531		540					
TAT	GCC	TAA	GTA	TAT	TTT	CCC	TAT	GTC	TAG	TAA	TAA	CCA
Tyr	Ala	.	Val	Tyr	Phe	Pro	Tyr	Val	.	.	.	Pro

549		558		567		576		585				
GAG	TCT	GTG	TAA	GCC	TGT	CAA	ATA	ACT	AAC	TCC	CCT	GTC
Glu	Ser	Val	.	Ala	Cys	Gln	Ile	Thr	Asn	Ser	Pro	Val

	594		603		612		621					
CCT	AGG	GTG	AAA	TGT	TAC	TCT	AAT	AAC	GTT	GGA	GAT	TTG
Pro	Arg	Val	Lys	Cys	Tyr	Ser	Asn	Asn	Val	Gly	Asp	Leu

Figure 2 A (cont. 2)

630	639	648	657	
CAT TCT GTG TTG TTT GTA GTA AGT TAT GGC TAG TAA TCT				
His Ser Val Leu Phe Val Val Ser Tyr Gly . . Ser				
666	675	684	693	702
ATT TAA GGT GAC TTG GAA TAC ATA AAA AAA AAA AAA AAA				
Ile . Gly Asp Leu Glu Tyr Ile Lys Lys Lys Lys Lys				
711	720	729	738	
AAA AAA AAA AAA AAA AAA AAA AAA AAA AAA AAA AAA AAA				
Lys Lys Lys Lys Lys Lys Lys Lys Lys Lys Lys Lys Lys Lys				
747				
AAA TGC A				
Lys Cys				

Figure 2 B

			11				20				29				38
CC	CTT	GTT	CTT	TTT	CAG	CTT	TCA	GTT	TTA	TTA	CTT	AGC			
	Leu	Val	Leu	Phe	Gln	Leu	Ser	Val	Leu	Leu	Leu	Ser			
			47				56				65				74
TCA	TTC	ACA	GTA	GTT	CTT	AGC	CAG	GAG	GAA	GAC	ATT	GGG			
Ser	Phe	Thr	Val	Val	Leu	Ser	Gln	Glu	Glu	Asp	Ile	Gly			
			83				92				101				110
GGT	TGG	TTT	ACC	ACC	AAA	CAT	CAT	GAC	CAC	CTT	TCA	CCA			
Gly	Trp	Phe	Thr	Thr	Lys	His	His	Asp	His	Leu	Ser	Pro			
			119				128				137				146
GCT	CAA	GCT	CCT	AAG	CCT	CAC	AAA	GGC	CAC	CAC	CAC	CCC			
Ala	Gln	Ala	Pro	Lys	Pro	His	Lys	Gly	His	His	His	Pro			
			164				173				182				191
AAA	CAT	TCC	CCA	GCC	CCT	TCA	CCA	ACT	AAG	CCT	CCC	ACT			
Lys	His	Ser	Pro	Ala	Pro	Ser	Pro	Thr	Lys	Pro	Pro	Thr			
			200				209				218				227
TAT	AGC	CCA	TCG	AAA	CCA	CCA	GTT	AAA	CCA	CCG	GTT	AAA			
Tyr	Ser	Pro	Ser	Lys	Pro	Pro	Val	Lys	Pro	Pro	Val	Lys			
			236				245				254				263
CCA	CCA	ACT	AAG	GCT	CCC	ACT	TAT	AGC	CCA	TCA	AAA	CCA			
Pro	Pro	Thr	Lys	Ala	Pro	Thr	Tyr	Ser	Pro	Ser	Lys	Pro			
			281				290				299				308
CCA	GCT	AAG	CCA	CCA	GTT	AAA	CCA	CCA	ACA	CCA	ACA	CCA			
Pro	Ala	Lys	Pro	Pro	Val	Lys	Pro	Pro	Thr	Pro	Thr	Pro			

Figure 2 B (cont. 1)

317	326	335	344	
TCA CCT TAT CCT GCT CCT GCT CCT ATT ACT AGG AAA CCT				
Ser Pro Tyr Pro Ala Pro Ala Pro Ile Thr Arg Lys Pro				
353	362	371	380	389
GTA GCA GTC CGT GGC CTT GTT TAC TGC AAG CCG TGC AAG				
Val Ala Val Arg Gly Leu Val Tyr Cys Lys Pro Cys Lys				
398	407	416	425	
TTT AGA GGG GTT AAA ACT CTA AAC CAA GCT TCC CCA CTC				
Phe Arg Gly Val Lys Thr Leu Asn Gln Ala Ser Pro Leu				
434	443	452	461	
CTG GGT GCG GTA GTG AAG CTA GTA TGC AAC AAC ACA AAG				
Leu Gly Ala Val Val Lys Leu Val Cys Asn Asn Thr Lys				
470	479	488	497	506
AAG ACA TTA GTG GAA CAG GGC AAG ACA GAC AAG AAT GGC				
Lys Thr Leu Val Glu Gln Gly Lys Thr Asp Lys Asn Gly				
515	524	533	542	
TTC TTC TGG ATC ATG CCC AAA TTC TTG TCC TCA GCA GCT				
Phe Phe Trp Ile MET Pro Lys Phe Leu Ser Ser Ala Ala				
551	560	569	578	
TAC CAC AAA TGC AAG GTG TTC TTG GTC TCA TCA AAC AAT				
Tyr His Lys Cys Lys Val Phe Leu Val Ser Ser Asn Asn				
587	596	605	614	623
ACT TAC TGT GAT GTC CCA ACA GAT TAC AAT GGT GGA AAA				
Thr Tyr Cys Asp Val Pro Thr Asp Tyr Asn Gly Gly Lys				

Figure 2 B (cont. 2)

	632		641		650		659
TCT	GGT	GCT	TTG	TTG	AAA	TAC	ACC
Ser	Gly	Ala	Leu	Leu	Lys	Tyr	Thr
	668		677		686		695
CCA	GCA	GCT	ACT	TCT	CTC	CCT	GTT
Pro	Ala	Ala	Thr	Ser	Leu	Pro	Val
704		713		722		731	
GAT	GTC	TTC	ACT	GTT	GGA	CCT	TTT
Asp	Val	Phe	Thr	Val	Gly	Pro	Phe
	749		758		767		776
AAG	AAG	GTG	CCA	TGC	AAA	AAG	TAA
Lys	Lys	Val	Pro	Cys	Lys	Lys	.
	785		794		803		812
AGA	AAG	ATA	GGA	AGG	AAA	AAT	TAA
Arg	Lys	Ile	Gly	Arg	Lys	Asn	.
821		830		839		848	
AGA	CGA	TTA	TGT	ACC	TGT	TTC	CTG
Arg	Arg	Leu	Cys	Thr	Cys	Phe	Leu
	866		875		884		893
TTT	ATT	AAT	AAA	TGA	AGC	AAA	GAG
Phe	Ile	Asn	Lys	.	Ser	Lys	Glu
	902		911		920		929
TCT	TGT	TTT	CCT	ATT	TTG	TTT	CTC
Ser	Cys	Phe	Pro	Ile	Leu	Phe	Leu

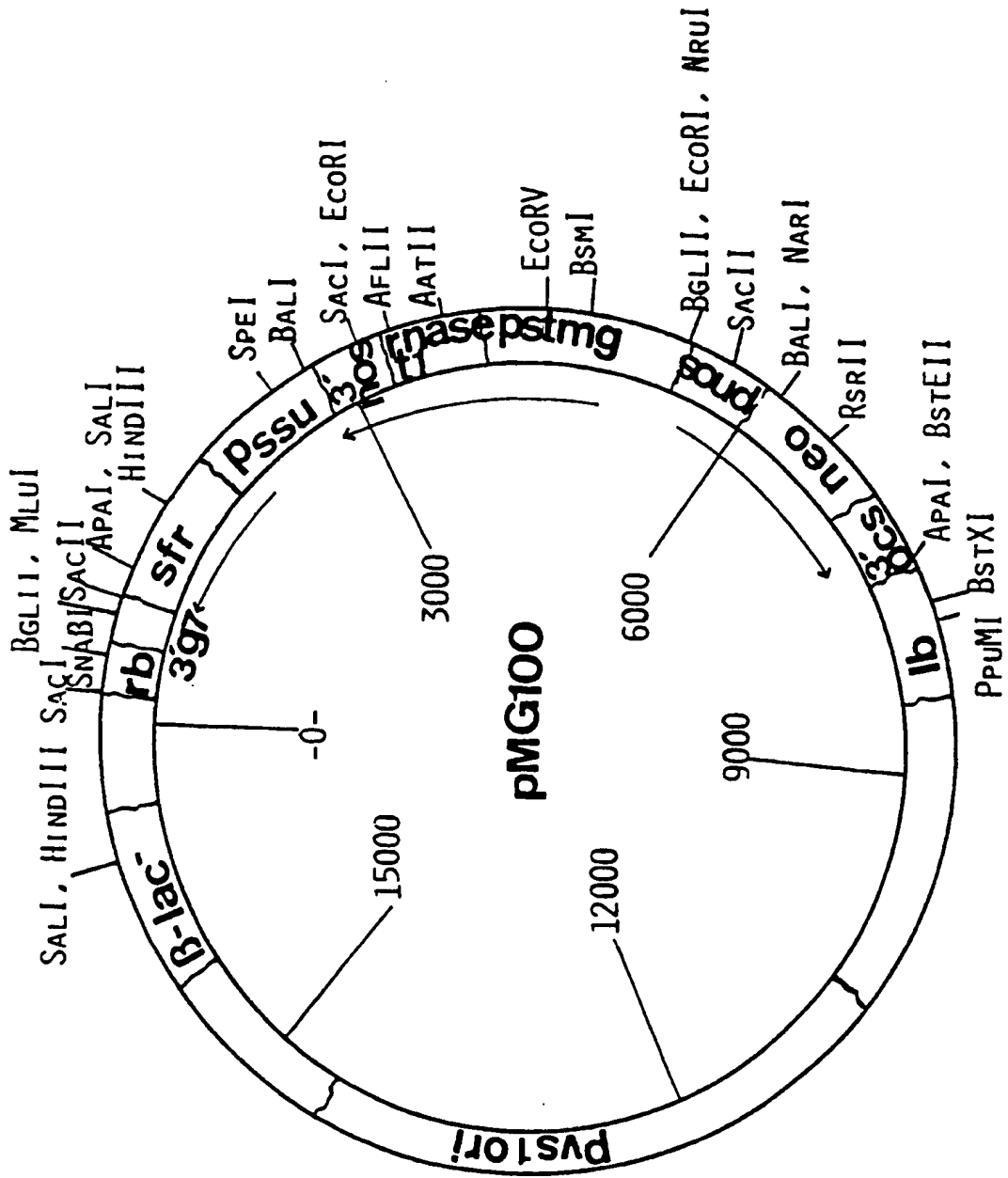
Figure 2 B (cont. 3)

938 947 956 965 974
CAA GTA AAA TGG ATT TAT AAG TTT TTC TTC AAA AAA AAA
Gln Val Lys Trp Ile Tyr Lys Phe Phe Phe Lys Lys Lys

983 992 1001 1010
AAA AAA AAA AAA AAA AAA AAA AAA AAA AAA AAA AAA
Lys Lys Lys Lys Lys Lys Lys Lys Lys Lys Lys Lys Lys

1019	1028	1037	1046
AAA AAA AAA AAA	AAA AAA AAA AAA	AAA AAT GCA GGT	CGA
Lys Lys Lys Lys	Lys Lys Lys Lys	Lys Asn Ala Gly	Arg

FIG.3



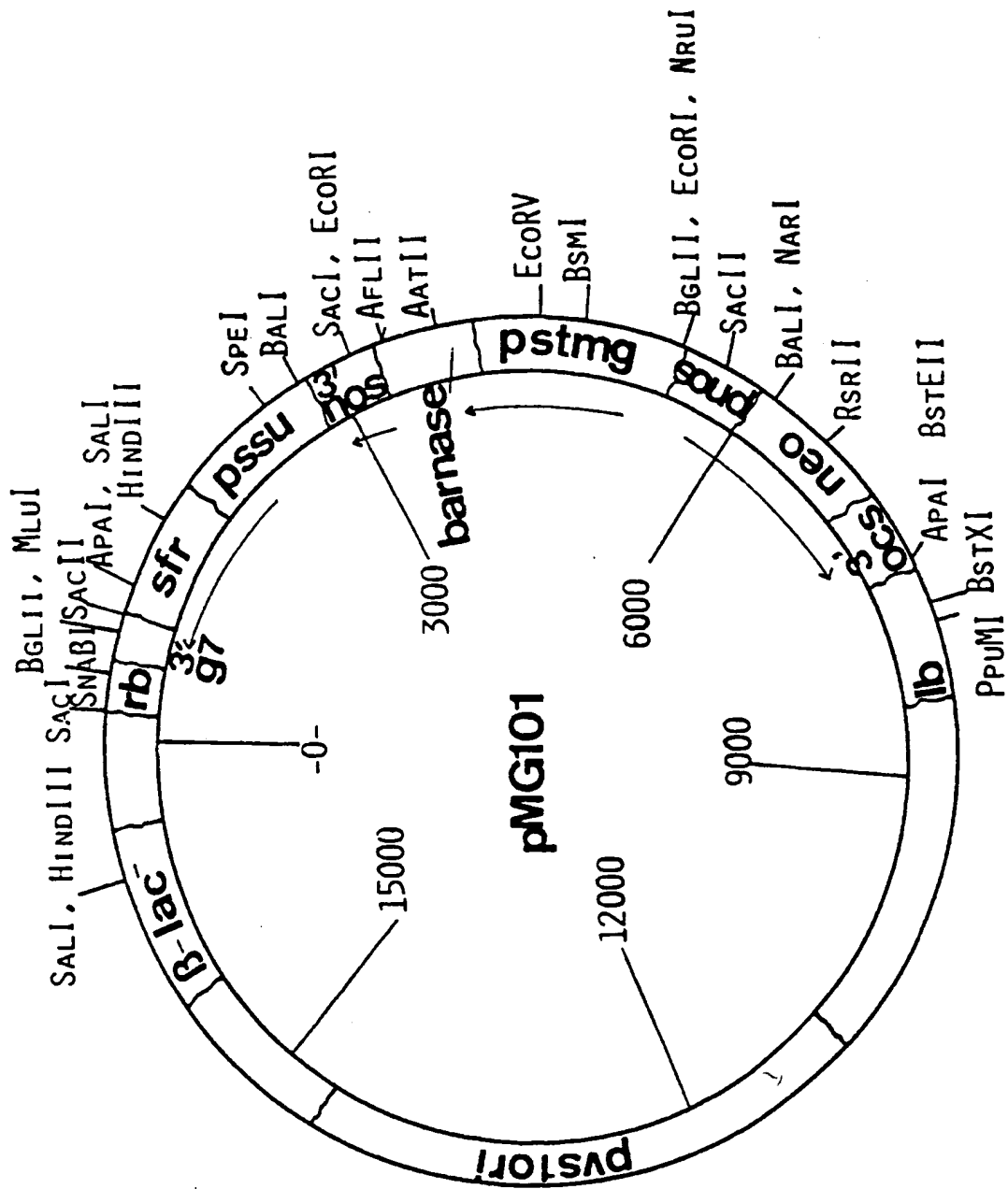


FIG. 4

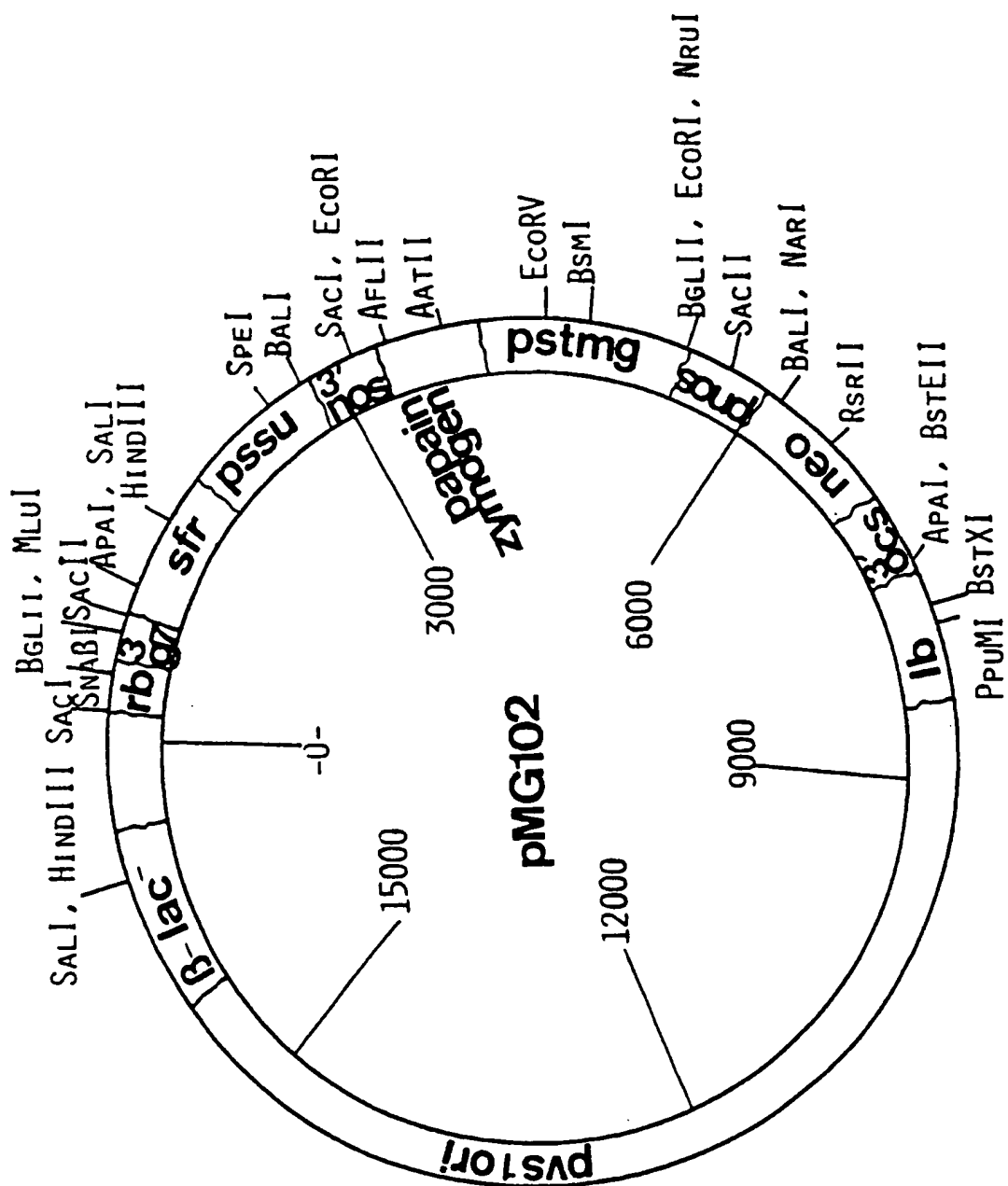
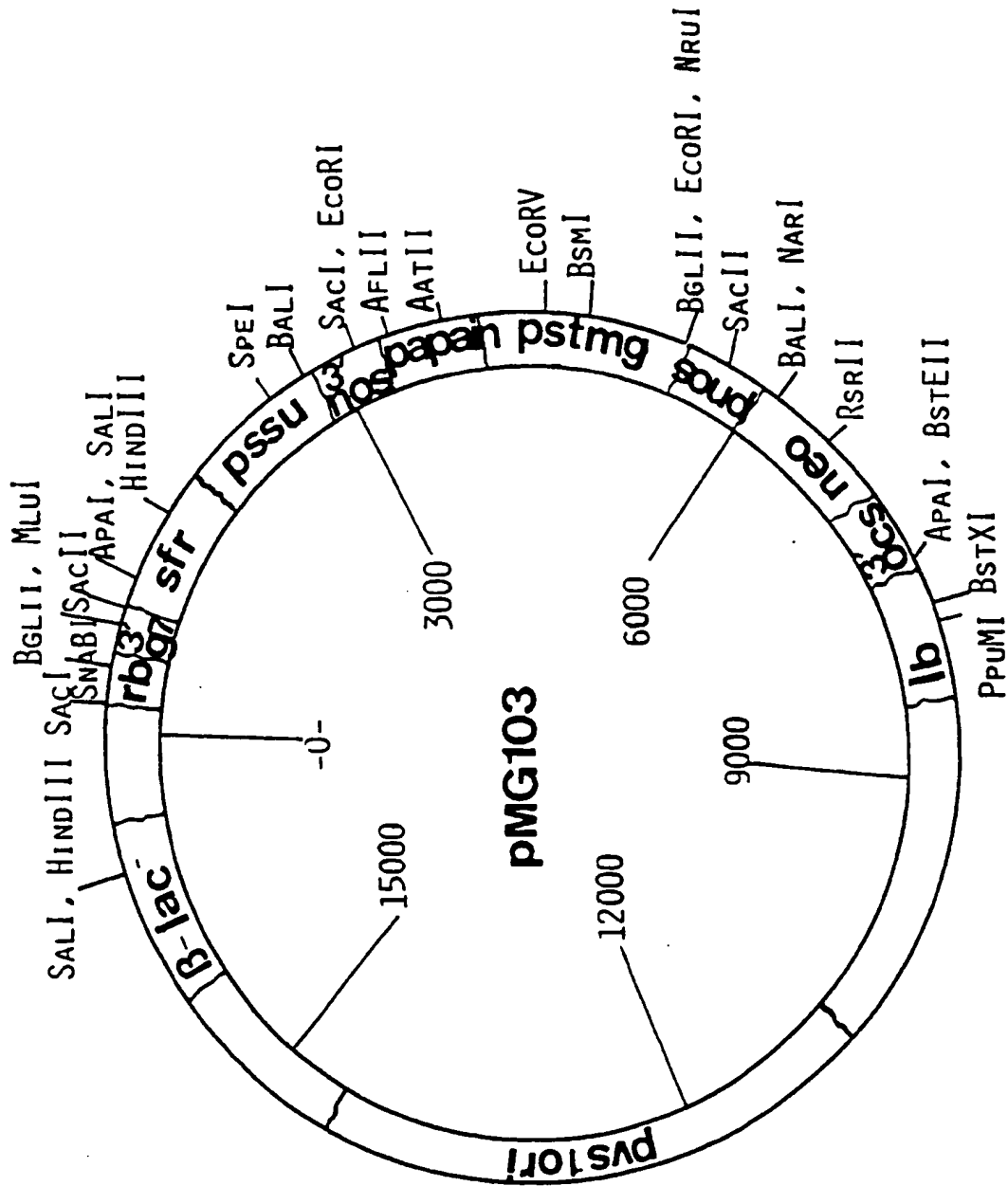


FIG. 5

FIG. 6



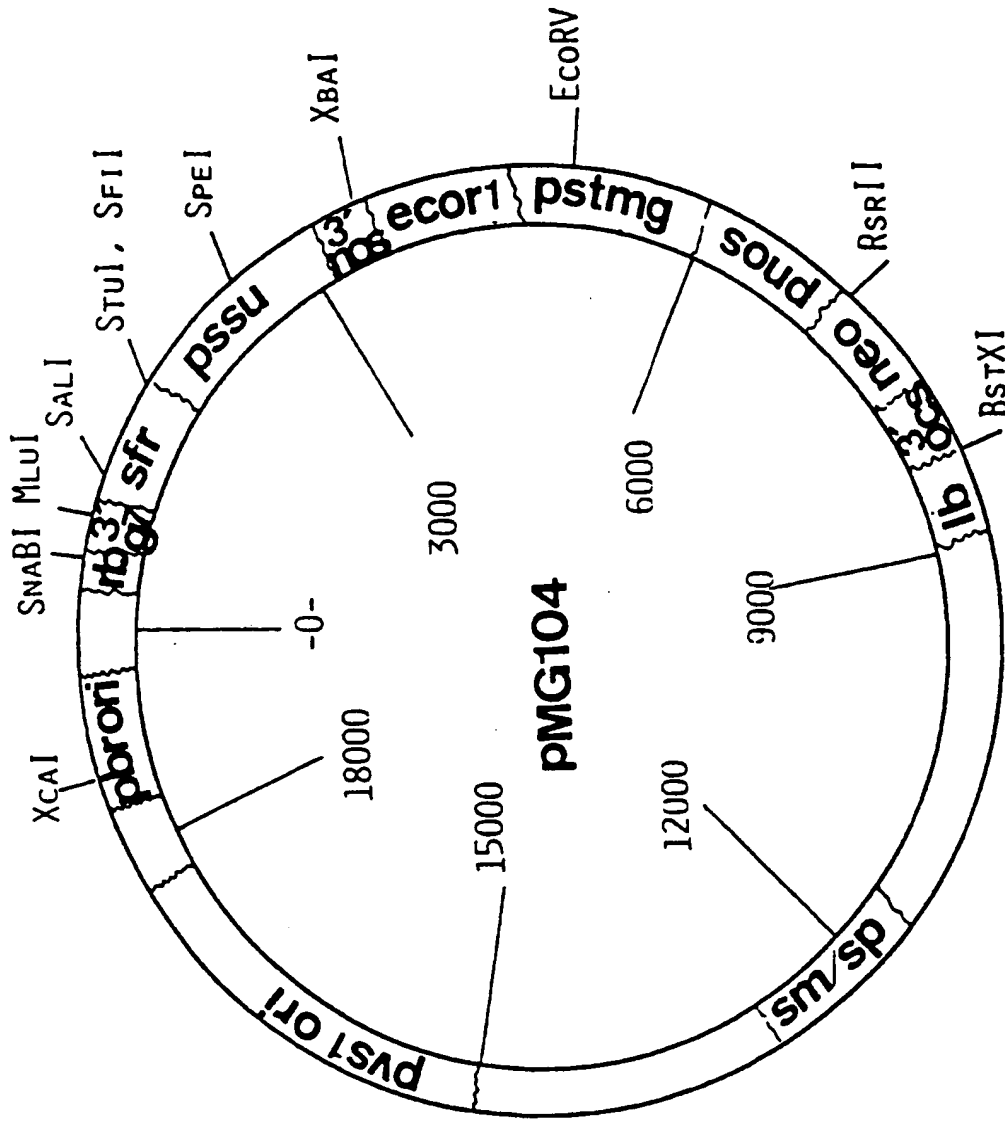


FIG. 7

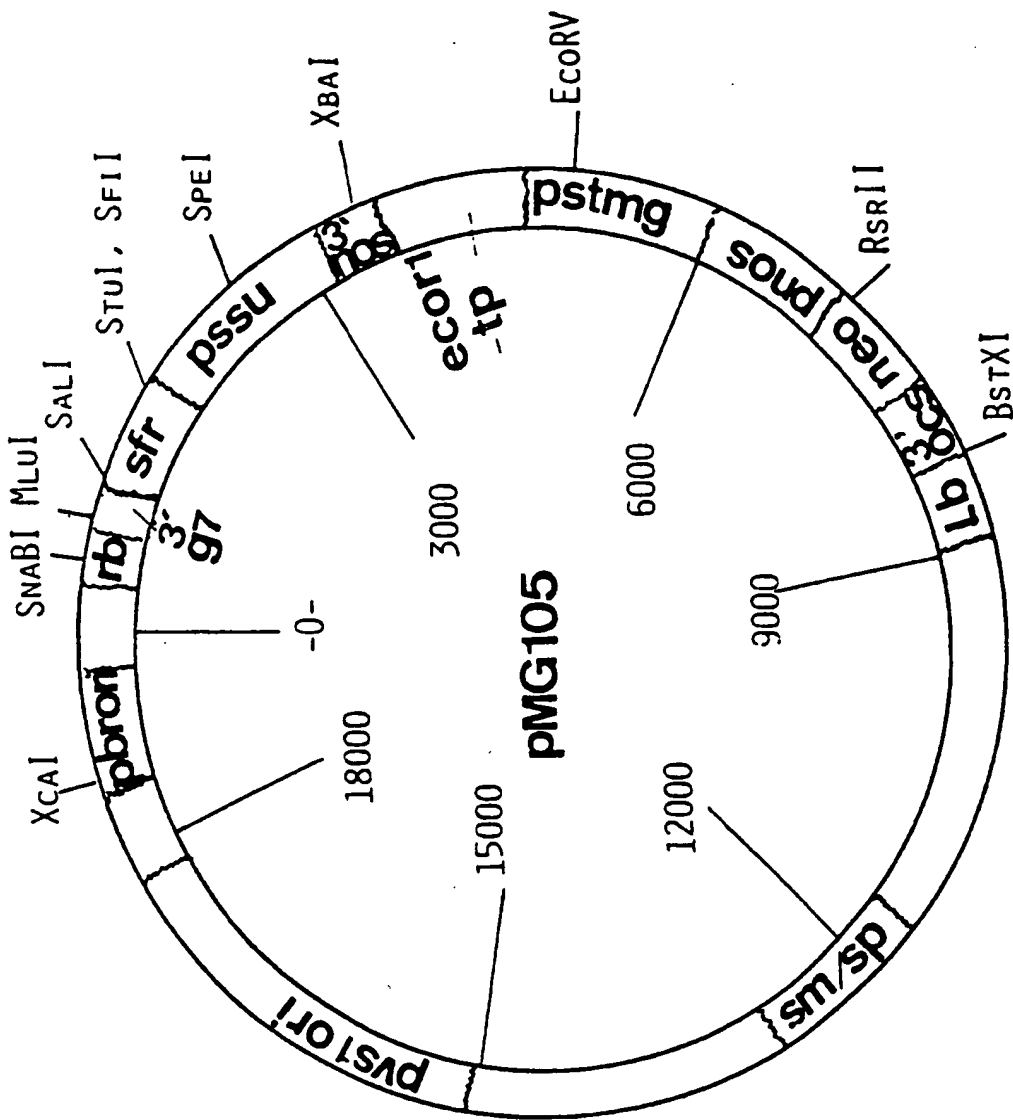


FIG. 8